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(54) Title: ALKYLATING AGENT COMBINATIONS IN THE TREATMENT OF CANCER

(57) Abstract: This application provides compositions and methods useful in the treatment of certain cancers. In part, this application is based on the recognition that certain molecules that target abasic lesions or AP sites in DNA improve, augment, or potentiate the chemotherapeutic efficacy of certain anticancer agents.

Alkylating Agent Combinations In the Treatment of Cancer

This invention was made in part with government support under grants
5 RO1CA73062, RO1ES06288, U01CA75525 and P30CA43703 from the USPHS,
and grant NIH-CA-86357 as well as a RAID initiative through the NCI as NSC3801.
The government has certain rights in the invention.

FIELD OF THE INVENTION

This invention generally relates to novel compositions and methods for the
10 treatment of certain cancers. Additionally, this invention relates to novel
compositions and methods to screen drugs useful for the treatment of certain
cancers.

BACKGROUND

Cancer is a worldwide problem. The American Cancer Society estimates that
15 over one half million people will die from cancer in the United States alone in 1999.
As such, finding novel compositions and methods for the treatment of cancer is of
vital interest. The treatment of cancer falls into three general categories:
chemotherapy, radiation therapy and surgery. Often, therapies are combined since a
combination of therapies often increases the probability the cancer will be eradicated
20 as compared to treatment strategies utilizing a single therapy. Most typically, the
surgical excision of large tumor masses is followed by chemotherapy and/or
radiation therapy.

Chemotherapeutic agents can work in a number of ways. For example,
chemotherapeutic can work by interfering with cell cycle progression or by
25 generating DNA strand breaks. If the cancer cell is not able to overcome the cell
cycle blockage or cell injury caused by the therapeutic compound, the cell will often
die via apoptotic mechanisms. The use of a single chemotherapeutic agent in the
treatment of cancer, with or without surgery or radiation, has several disadvantages.
First, the cells may develop resistance to the chemotherapeutic agent. Such
30 resistance results either in the requirement for higher dosages of the drug and/or the
renewed spread of the cancer. Chemotherapeutic agents can be toxic to the patient.

Therefore, there is a practical upper limit to the amount that a patient can receive. However, if two chemotherapeutic agents are used in concert, the dosage of any single drug can be lowered. This is beneficial to the patient since using lower levels of chemotherapeutic agents is generally safer for the patient. Additionally, cancer
5 cells are less likely to generate resistance to the combination of drugs as they are to a single drug.

The design of drug combinations for the chemotherapeutic treatment of cancer should be approached with the goals of 1) finding a combination that is synergistic with and not merely additive to the first compound with respect to the
10 elimination of the tumor, and 2) finding a second drug that does not potentiate the toxic effects of the first chemotherapeutic agent. These conditions require a great deal of empirical testing of agents known to have anticancer properties with agents that either may have anticancer properties, or that may augment the first agent in other ways. TMZ is currently employed in chemotherapeutic treatment of certain
15 tumors. It works by dramatically increasing the mutation rate of cells undergoing DNA replication. Such cells, because of the high number of mutations which they have acquired as a result of the treatment with TMZ, are rapidly removed by apoptosis, thereby potentially eliminating the tumor. Some tumor cells are resistant to treatment by TMZ due to deficiencies in the mismatch repair (MMR) system in
20 the cell. A defective MMR system prevents the cell from recognizing O⁶mG DNA adducts thereby making the cell resistant to elimination.

Baer *et al.*, in U.S. Pat.No. 5,731,304, note that the toxicity of temozolomide can be potentiated by agents that inhibit the enzyme O⁶-alkylguanine DNA
alkyltransferase (ATase). In particular, they note that O⁶-benzylguanine (BG) can
25 enhance the toxicity of temozolomide in certain cell lines that exhibit high levels of ATase (*e.g.* 300-fold in MAWI cells). However, in other cell lines that exhibit lower levels of ATase (*e.g.* U373 cells) the effect is significantly less.

Mitchell and Dolan (*Cancer Chemother Pharmacol* 32:59-63, 1993) note that temozolomide (TMZ) and an analogue, 5-(dimethyltriazeno)imidazole-4-
30 caroxamide (DITC), can be effective in enhancing the anti-tumor effects of 1,3-bis(2-chloroethyl) 1-nitrosourea (BCNU). TMZ and DITC work by depleting cells or tumors of O⁶-alkylguanine-DNA alkyltransferase (AGT). AGT is a DNA repair

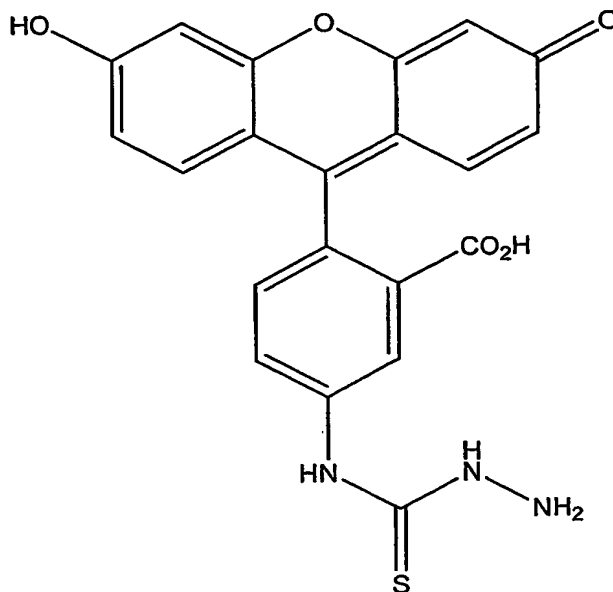
protein that selectively removes adducts from the O⁶ position of guanine in DNA by a stoichiometric transfer of the alkyl group to a cysteine moiety. Removal of the alkyl group from the DNA by methylation of the O⁶ position of the guanine effectively inactivates the AGT. As with the patent referred to above, the disclosed
5 method is limited to specific cells or cancers.

Therefore, what is needed are therapies that utilize the synergistic properties of two or more therapeutic agents for the treatment of cancer that have a broader range of targets or a different range of targets than those combination therapies already known.

10 SUMMARY OF THE INVENTION

This application provides compositions and methods useful in the treatment of certain cancers. In part, this application is based on the recognition that certain molecules that target abasic lesions or AP sites in DNA improve, augment, or potentiate the chemotherapeutic efficacy of certain anticancer agents.

15 The present invention relates generally to compositions and methods for the treatment of hyperproliferation, such as cancer, that utilize both a chemotherapeutic that induces the formation of AP sites in the DNA of a cell and an inhibitor of the base excision repair pathway (BER pathway). In preferred embodiments, the BER inhibitor is methoxyamine or compound A.



Compound A

In one embodiment, this application provides methods for enhancing the therapeutic effect of an anticancer agent, such as, for example, an anticancer agent that induces the formation of AP sites, by administering a base excision repair (BER) inhibitor, whereby the BER inhibitor enhances the effect of the anticancer agent. The BER inhibitor may be administered either prior to the administration of the anticancer agent or at the same time. It may be administered either orally or intravenously.

The certain embodiments, the BER inhibitor may be any one of an AP endonuclease inhibitor, a DNA glycosylase inhibitor, a DNA polymerase inhibitor, a PARP inhibitor, a DNA alkyltransferase inhibitor, and a DNA ligase inhibitor.

In one aspect, the AP endonuclease inhibitor may be anyone of methoxyamine, compound A, and N-ethylmaleimide. In other aspects, the BER inhibitor is a PARP inhibitor selected from PD128763, 3-AB, 6-AN, NU1025, and 4-amino-1,8-naphthalimide. In another aspect the BER inhibitor is an inhibitor of DNA polymerase, such as DNA polymerase β , γ , or ϵ . The DNA polymerase inhibitor may be selected from prunasin; aphidicolin; ddCTP; ddTTP; ddATP; ddGTP; Ara-C; and arabinocytidine.

In yet other aspects, the BER inhibitor is a DNA ligase inhibitor that inhibits the action of DNA ligase I or DNA ligase II. In other embodiments, the BER inhibitor is a DNA glycosylase inhibitor.

In certain aspects of the invention, the anticancer agent is an agent that induces the formation of AP sites and may be selected from any one of the following agents: a DNA oxidizing agent; ultraviolet radiation; a DNA intercalating agent; a radiosensitizing agent; a cross-linking agent; and an alkylating agent.

In certain embodiments, examples of the alkylating agents include TMZ (temozolomide); MeOSO₂(CH₂)₂-lextropsin (Me-Lex); Cis-diamminedichloroplatinum II (cis-DDP); mitomycin bioreductive alkylating agents; quinones; STZ (streptozotocin); cyclophosphamide; nitrogen mustard family members such as chloroambucil; pentostatin (purine analogs); fludarabine; bendamustine hydrochloride which is the active ingredient of Ribomustin (alkylating group in common with the nitrogen mustard family, also an antimetabolites);

chloroethylating nitrosoureas (lomustine, fotemustine, cystemustine); dacarbazine (DTIC); and procarbazine.

In other aspects, the anticancer agent is a DNA oxidizing agent and the DNA oxidizing agent may be any one of the following: hydrogen peroxide, bleomycin, and adriamycin. In yet other aspects, the agents that induce the formation of AP sites include ultraviolet radiation, gamma irradiation, UVA, UVB, and radiosensitizing agent. Examples of radiosensitizing agents include agents such as IUdR.

In yet other aspects, agents that induce the formation of AP sites include cross-linking agents such as BCNU; intercalating agents such as bleomycin; adriamycin; quinacrine; echinomycin; and anthrapyrazoles. In other embodiments, the anticancer agent may be any one of uracil; hypoxanthine; 5-FU; flavonoids; ecteinascidin-743; cis-diamminedichloroplatinum II; 6-thioguanine; taxol; and camptothecin analogs.

The formulations and compositions disclosed herein may be useful in the treatment of cancers such as carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.

In yet another embodiment, the application provides kits comprising: 1) a first pharmaceutical preparation comprising an anticancer agent that induces formation of AP sites; and 2) a second pharmaceutical preparation comprising a base excision repair inhibitor, and instructions for administering the first and second pharmaceutical preparations to a patient for the treatment of cancer.

In certain embodiments, the invention contemplates that anticancer agents such as temozolomide (TMZ) may be used together with BER inhibitors such as methoxyamine (MX) as a treatment for certain tumors that are resistant to treatment by the anticancer agent alone. Additionally, the invention contemplates that anticancer agents such as TMZ and inhibitors of poly-ADP-ribose polymerase (PARP) may be used as a treatment for certain tumors that are resistant to treatment by the anticancer agent alone. The present invention generally comprises novel compositions to 1) screen for compounds that can potentiate or modulate the therapeutic effect of anticancer agents such as temozolomide (TMZ), either alone or

combined with methoxyamine (MX), 2) provide model systems for the study of cancer treatments by agents that modulate DNA repair mechanisms and 3) provide treatments for certain hyperproliferative disorders such as cancers.

With regard to the treatment of cancer, the present invention contemplates methods of treating cancer that utilize an anticancer agent such as TMZ in
5 conjunction with another agent that is capable of potentiating the toxic effect of the anticancer agent. More specifically, the invention relates to potentiating the effect of anticancer agents such as TMZ with agents that modify DNA adducts created by the anticancer agent. Even more specifically, the invention relates to potentiating the
10 effect of anticancer agents such as TMZ with agents that interact with AP sites and/or modify N⁷-methylguanine (N⁷mG) and N³-methyladenine (N³mA) DNA adducts. Examples of such agents are MX and compound A. The present invention is not limited by the order in which the agents are administered. In one embodiment, the agents are administered sequentially. In another embodiment, the agents are
15 administered as a combined formulation (e.g., a formulation comprising methoxyamine and temozolomide). In one embodiment, the present invention contemplates a method of treating cancer by administering to a patient diagnosed with cancer a first formulation comprising a BER inhibitor such as methoxyamine and a second formulation comprising an anticancer agent such as temozolomide. In
20 certain alternative embodiments, the anticancer agent and the BER inhibitor are administered to the patient in a single formulation.

Additionally, the invention relates to potentiating anticancer agents such as temozolomide with agents that interfere with the ability of PARP to be effective in DNA repair by either competing with PARP, inhibiting PARP, or degrading PARP.
25 The present invention is not limited to the agent used to inhibit PARP activity. In one embodiment, PD128763 is used. In another embodiment 6-AN is used. In another embodiment, 3-AB is used. The present invention is not limited by the order in which the agents are administered. In one embodiment the agents are administered sequentially. In another embodiment, the agents are administered as a
30 combined formulation. In one embodiment, the present invention contemplates a method of treating cancer by administering to a patient diagnosed with cancer a first formulation comprising a poly-(ADP-ribose)-polymerase inhibitor and a second

formulation comprising an anticancer agent such as temozolomide. In certain alternative embodiments, the inhibitor and the agent are administered to the patient in a single formulation.

The present invention is not limited to the method of administration of the treatment. In one embodiment, the treatment is administered orally. In another embodiment the treatment is administered intravenously. In yet another embodiment, the treatment is administered intraperitoneally. In yet another embodiment, the treatment is administered directly to the tumor by injection or, in the case of skin tumors, for example, by direct application of creams or ointments. In certain embodiments, one agent is administered by one route, while the second agent is administered by a second route (e.g., temozolomide is administered orally, while the BER inhibitor is administered by injection).

As noted above, the present invention also contemplates screening assays to identify drugs that augment the ability of anticancer agents such as temozolomide to inhibit tumor growth, drugs that augment the ability of temozolomide with methoxyamine to inhibit tumor growth, and drugs that augment the ability of temozolomide with PARP inhibitors to inhibit tumor growth. A variety of assay formats are contemplated for testing the potential of compounds suspected of augmenting the anti-tumor effect of anticancer agents such as TMZ. In one embodiment, cells are pretreated with the test compound that may augment the antitumor effect of the agent, followed by treatment with the anticancer agent. Cell growth and/or cell death are then measured to determine if there is an anti-tumor effect. In another embodiment, cells are treated with the test compound that may augment the anti-tumor effect of the agent at approximately the same time as they are treated with the anticancer agent. Cell growth and/or cell death are then measured to determine if the regime had an anti-tumor effect. The invention is not limited to any particular measurement technique for apoptosis or cell growth. Various methods are envisioned. For example, mitosis can be measured by use of fluorescent dyes that intercalate into DNA, by the measurement of ³H-thymidine incorporation, or by colorimetric assays. Such assays permit the use of high-throughput screening methods. Apoptosis can be measured by the use of fluorescent

dyes that intercalate into DNA, annexin-V staining of phosphatidyl serine residues of the cell surface or morphological changes in cell appearance.

The present invention contemplates a method of screening for compounds that augment the anti-tumor effect of an anticancer agent such as temozolomide by contacting tumor cells (e.g., in vitro, or optionally in vivo) with a first formulation comprising a test compound that may potentiate the anticancer agent and a second formulation comprising the anticancer agent and measuring the growth and death rates of said cells to determine whether the test compound potentiates the activity of the anticancer agent.

The present invention contemplates a method of screening for compounds that augment the anti-tumor effect of an anticancer agent such as temozolomide by implanting tumor cells in an animal, and administering to the animal a first formulation comprising a test compound that may potentiate an anticancer agent such as temozolomide and a second formulation comprising the anticancer agent, and measuring the growth and death rates of said cancer cells to determine whether the test compounds potentiates the anticancer agent.

It is not intended that the present invention be limited by the nature of the compounds to be screened in the screening assay of the present invention. For example, a variety of compounds including peptides, organic compounds, and inorganic compounds, are contemplated. Additionally, combinations of compounds are contemplated by the present invention.

In some embodiments, a method is provided in which a patient diagnosed with cancer is administered a dose of a first formulation comprising a BER inhibitor such as methoxyamine and a dose of a second formulation comprising an anticancer agent such as 1,3-bis (chloroethyl) 2-nitrosourea (BCNU), wherein said inhibitor is administered in an amount sufficient to potentiate toxicity of said agent. In some embodiments, the inhibitor and agent are administered sequentially, while in other embodiments the inhibitor and agent are administered in a single formulation. In some embodiments, said the inhibitor and agent are administered orally, while in other embodiments, said the inhibitor and agent are administered intravenously. In yet other embodiments, a formulation comprising both the inhibitor and agent is contemplated.

In other embodiments, a method is provided comprising administering to a patient diagnosed with cancer a first formulation comprising a BER inhibitor such as methoxyamine and a second formulation comprising an anticancer drug or agent that exerts cytotoxicity mediated by oxidative DNA damage, wherein said BER inhibitor
5 is administered in an amount sufficient to potentiate toxicity of said anticancer agent or drug. In some embodiments, said anticancer drug or agent is selected from bleomycin and adriamycin. In some embodiments, said inhibitor and said anticancer drug or agent are administered sequentially. In other embodiments, said inhibitor and said anticancer drug or agent are administered as a formulation.

10 In other embodiments, a method is provided comprising administering to a patient diagnosed with cancer a first formulation comprising a BER inhibitor such as methoxyamine and a second formulation comprising an anticancer drug or agent selected from hypoxanthine, 5-FU, uracil, IUdR, bleomycin and adriamycin, wherein said inhibitor is administered in an amount sufficient to potentiate toxicity
15 of said anticancer drug or agent. In some embodiments, said inhibitor and said anticancer drug or agent are administered sequentially, while in other embodiments, said inhibitor and said anticancer drug or agent are administered in a single formulation. In some embodiments, said administration is oral administration, in other embodiments, said administration is intravenous administration.

20 In some embodiments, a formulation comprising a BER inhibitor such as methoxyamine and an anticancer drug or agent selected from hypoxanthine, 5-FU, uracil, IUdR, bleomycin and adriamycin is contemplated. In some embodiments, said anticancer drug or agent is BAR. In some embodiments, a formulation comprising methoxyamine and hydrogen peroxide is contemplated.

25 In some embodiments, a method is provided comprising administering to a patient diagnosed with cancer a formulation comprising a BER inhibitor such as methoxyamine and treating said patient with radiation; wherein said inhibitor is administered in an amount sufficient to potentiate the toxicity of said radiation. In some embodiments, said inhibitor administration and radiation treatment occur
30 sequentially, while in other embodiments, said inhibitor administration occurs essentially simultaneously with said radiation treatment.

In some embodiments, a method is provided comprising administering to a patient diagnosed with cancer a first formulation comprising a BER inhibitor such as methoxyamine and a second formulation comprising a DNA oxidizing agent such as hydrogen peroxide, wherein said inhibitor is administered in an amount sufficient to
5 potentiate the cytotoxic effects of said oxidizing agent.

In other embodiments, a method is provided comprising administering to a patient diagnosed with cancer a first formulation comprising a BER inhibitor such as methoxyamine and a second formulation comprising an antineoplastic agent such as iododeoxyuridine (IUdR), wherein said methoxyamine is administered in an amount
10 sufficient to further increase the radiosensitivity of the tumor cells in said patient. In some embodiments, the method further comprises treating said patient with radiation therapy. In some embodiments, said the inhibitor and agent are administered sequentially, while in other embodiments, they are administered as a formulation. In some embodiments, the inhibitor and agent are administered orally, while in other
15 embodiments, said they are administered intravenously.

In certain embodiments, the compositions and methods of the invention are used to treat a patient having a cancer selected from carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.

20

DESCRIPTION OF THE FIGURES

Figure 1. MX increases the sensitivity of colon cancer cells to TMZ. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 6 mM MX or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. A. HTC116 cells; B. SW480 cells. (■),
25 TMZ alone; (◆), MX plus TMZ; (●), BG plus TMZ; (▲), BG and MX plus TMZ. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 2. Cytotoxicity of MMS enhanced by MX in colon cancer cell lines. Cells were treated with 0-3 mM MMS for 1 h or MMS and 6 mM MX plus 10 μ M

BG for 2 h prior to 2 h exposure to MMS. A. HCT116 cells; B. SW480 cells. (■), MMS alone; (◆), MX plus MMS; (●), BG plus MMS; (▲), BG and MX plus MMS. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

5 Figure 3. Inhibitors of PARP enhance cytotoxicity of TMZ in MMR wt SW480 cells. A. Cells were treated with 0-1500 TMZ for 2 h or TMZ and 100 μ M PD128763 or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (◆), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus PD128763. B. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 10 μ M 6-AN (pretreated for 48 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (◆), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN. C. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 3 mM 3-AB (pretreated for 24 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (▲), BG plus TMZ; (●), 3-AB plus TMZ; (▲), BG and TMZ plus 3-AB. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

10 Figure 4. Inhibitors of PARP enhance cytotoxicity of TMZ in MMR deficient HCT116 cells. A. Cells were treated with 0-1500 TMZ for 2 h or TMZ and 100 μ M PD128763 or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (▲), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus PD128763. B. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 100 μ M 6-AN (pretreated for 48 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (▲), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN. C. Cells were treated with 0-1500 μ M TMZ for 2 h or TMZ and 3 mM 3-AB (pretreated for 24 h) or plus 10 μ M BG for 2 h prior to 2 h exposure to TMZ. (■), TMZ alone; (◆), BG plus TMZ; (●), 3-AB plus TMZ; (A), BG and TMZ plus 3-

AB. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 5. Synergy analysis of the interaction between TMZ and MX (A) or PD128763 (B) in HTC116 cells.

5 Figure 6. Survival fraction of human colon cancer cell lines after exposure to BCNU plus MX. Cells were treated with 0-100 μ M BCNU for 2 h or BCNU and 6 mM MX plus 10 μ M BG for 2 h prior to 2 h exposure to BCNU. A. HCT116 cells; B. SW480 cells. (■), BCNU alone; (◆), MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU. Results are the mean \pm SD of at least three separate
10 experiments performed in duplicate.

Figure 7. Survival fraction of human colon cancer cell lines after exposure to BCNU plus PD128763. Cells were treated with 0-100 μ M BCNU for 2 h or BCNU and 100 μ M PD 128763 plus 10 μ M BG for 2 h prior to 2 h exposure to BCNU. A. HCT116 cells; B. SW480 cells. (■), BNCU alone; (◆), PD 128763 plus BCNU;
15 (●) BG plus BCNU; (▲), BG and PD 128763 plus BCNU; (▶), BG, PD128763 and MX plus BCNU. Results are the mean \pm SD of at least three separate experiments performed in duplicate.

Figure 8. Distribution of cell cycle and apoptosis in colon cancer cell lines after treatment with TMZ plus modifiers. A. MMR wt SW480 cells; B. MMR-deficient HCT116 cells.
20

Figure 9. PARP cleavage in colon cancer cells treated with TMZ plus modifiers. A. Controls, as labeled. B. SW480 cells; C. HCT116 cells; Lane 1, untreated; 2, TMX plus MX; 3, TMZ plus MX plus BG; 4, TMZ plus 3-AB; 5, TMZ plus 3-AB plus BG; 6, TMZ plus 6-AN; 7, TMZ plus 6-AN plus BG; 8, TMZ plus
25 PD128763; 9, TMZ plus PD128763 plus BG.

Figure 10. The effect of MX plus TMZ on growth of SW480 xenograft.

Figure 11. The effect of MX plus TMZ on growth of HCT116 xenograft.

Figure 12A shows the effects of BCNU alone and BCNU in combination with MX on HCT116 tumors grown in nude mice. Figure 12B shows the effects of
30 BCNU alone and BCNU in combination with MX on HCT116-Ch3 tumors grown in

nude mice. Figure 12C shows the effects of BCNU alone and BCNU in combination with MX on SW480 tumors grown in nude mice.

Figure 13 shows the effects of hydrogen peroxide (H_2O_2) alone and H_2O_2 in combination with MX on HCT116 cells in a growth inhibition assay.

5 Figure 14 shows that Compound A and MX have the same reactivity to AP sites. (A) Binding of A or MX to AP sites as assayed by ARP. (B) APE cleavage assays in regular and modified AP sites.

Figure 15 shows that A potentiates TMZ cytotoxicity in two tumor cells: DLD1 cells and HCT116 cells.

10 Figure 16 shows enhanced inhibition of tumor growth by TMZ and compound A in nude mice.

Figure 17 shows a schematic diagram of the preparation of DNA substrates with regular AP sites or MX-AP sites.

15 Figure 18 shows that MX-bound AP sites are resistant to cleavage by AP-endonuclease (APE).

Figure 19 shows topo II-mediated DNA cleavage in oligonucleotides containing a topo II cleavage site, in the presence or absence of etoposide.

20 Figure 20 shows DNA double strand breaks (DSBs) induced by the combination of TMZ, MX, and etoposide, at 4 hours and 72 hours after treatment, respectively.

25 Figure 21 shows colony survival assays in the cell line SW480 after either individual treatment with TMZ (A or D) or etoposide (C), or combination treatment with TMZ and MX (A), TMZ, MX and etoposide (A), or TMZ and etoposide (D). The synergy analysis of the interaction among TMZ, MX, and etoposide is indicated by the combination index (CI) (shown in B and E).

Figure 22A shows that MX inhibited detection of AP sites by ARP; Figure 22B shows the relative reduction of AP sites estimated by densitometric intensity, indicating that MX (5-20 mM) blocked 38-80% of AP binding sites.

30 Figure 23A is a diagram depicting the preparation of DNA substrates containing either regular AP sites or MX-modified AP sites; Figure 23B shows that MX-AP sites are resistant to cleavage by APE.

DEFINITIONS

To facilitate understanding of the invention, a number of terms are defined below.

5 As used herein "agent" or "drug" is used herein to denote a chemical compound, a mixture of chemical compounds, a biological macromolecule, or an extract made from biological materials such as bacteria, plants, fungi, or animal (particularly mammalian) cells or tissues that are suspected of having therapeutic properties. The agent or drug may be purified, substantially purified or partially
10 purified.

As used herein "agonist" refers to molecules or compounds which mimic the action of a "native" or "natural" compound. Agonists may be homologous to these natural compounds in respect to conformation, charge or other characteristics. Thus, agonists may be recognized by, *e.g.*, receptors expressed on cell surfaces. This
15 recognition may result in physiologic and/or biochemical changes within the cell, such that the cell reacts to the presence of the agonist in the same manner as if the natural compound was present.

As used herein "antagonist" refers to molecules or compounds which inhibit the action of a "native" or "natural" compound. Antagonists may or may not be
20 homologous to these natural compounds in respect to conformation, charge or other characteristics. Thus, antagonists may be recognized by the same or different receptors or molecules that are recognized by an agonist. Antagonists may have allosteric effects which prevent the action of an agonist (*e.g.*, by modifying a DNA adduct, or antagonists may prevent the function of the agonist (*e.g.*, by blocking a
25 DNA repair molecule).

As used herein, the term "fusion protein" refers to a chimeric protein containing the protein of interest joined to an exogenous protein fragment. The fusion partner may provide a detectable moiety, may provide an affinity tag to allow purification of the recombinant fusion protein from the host cell, or both. If desired,
30 the fusion protein may be removed from the protein of interest by a variety of enzymatic or chemical means known to the art.

As used herein, the term "purified" or "to purify" refers to the removal of one or more contaminants from a sample. The present invention contemplates purified compositions (discussed above).

As used herein, the term "partially purified" refers to the removal of a moderate portion of the contaminants of a sample to the extent that the substance of interest is recognizable by techniques known to those skilled in the art as accounting for a measurable amount of the mixture. Preferably, the compound of interest is at least 5% of the total preparation and up to 50% of the total preparation.

As used herein, the term "substantially purified" refers to the removal of a significant portion of the contaminants of a sample to the extent that the substance of interest is recognizable by techniques known to those skilled in the art as the most abundant substance in the mixture.

As used herein the term "portion" when in reference to a protein (as in "a portion of a given protein") refers to fragments of that protein. The fragments may range in size from four amino acid residues to the entire amino acid sequence minus one amino acid. In one embodiment, the present invention contemplates "functional portions" of a protein. Such portions are "functional" if they contain a binding region (i.e., a region having affinity for another molecule) and such binding can take place (i.e., the binding region functions, albeit with perhaps lower affinity than that observed for the full-length protein). Such "functional portions" of the gene product are typically greater than 10 amino acids in length, and more typically greater than 50 amino acids in length, and even more typically greater than 100 amino acids in length. "Functional portions" may also be "conserved portions" of the protein. The alignment of the various gene products permit one skilled in the art to select conserved portions of the protein (i.e., those portions in common between two or more species) as well as unconserved portions (i.e., those portions unique to two or more species). In one embodiment, the present invention contemplates conserved portions 10 amino acids in length or greater, and more typically greater than 50 amino acids in length.

"Staining" shall be defined as any number of processes known to those in the field that are used to better visualize, distinguish or identify a specific component(s) and/or feature(s) of a cell or cells.

"Immunofluorescence" is a staining technique used to identify, mark, label, visualize or make readily apparent by procedures known to those practiced in the art, where a ligand (usually an antibody) is bound to a receptor (usually an antigen) and such ligand, if an antibody, is conjugated to a fluorescent molecule, or the ligand is then bound by an antibody specific for the ligand, and said antibody is conjugated to a fluorescent molecule, where said fluorescent molecule can be visualized with the appropriate instrument (*e.g.*, a fluorescent microscope).

"Antibody" shall be defined as a glycoprotein produced by B cells that binds with high specificity to the agent (usually, but not always, a peptide), or a structurally similar agent, that generated its production. Antibodies may be produced by any of the known methodologies [Current Protocols in Immunology (1998) John Wiley and Sons, Inc., N.Y.] and may be either polyclonal or monoclonal.

"In operable combination", "in operable order" and "operably linked" as used herein refer to the linkage of nucleic acid sequences in such a manner that a nucleic acid molecule capable of directing the transcription of a given gene and/or the synthesis of a desired protein molecule is produced. The term also refers to the linkage of amino acid sequences in such a manner so that a functional protein is produced.

"Antigen" shall be defined as a protein, glycoprotein, lipoprotein, lipid or other substance that is reactive with an antibody specific for a portion of the molecule.

"Morphology" shall be defined as the visual appearance of a cell or organism when viewed with the eye, a light microscope, a confocal microscope or an electron microscope, as appropriate.

"Patient" shall be defined as a human or other animal, such as farm animals or laboratory animals (*e.g.*, guinea pig or mice) capable of having cell cycle (influenced) determined diseases, either naturally occurring or induced, including but not limited to cancer.

A "wild type" (wt) cell or cell line shall be defined as a cell or cell line that retains the characteristics normally associated with that type of cell or cell line for the physiological process or morphological characteristic that is being examined. It is permissible for the cell or cell line to have non-wild type characteristics for

physiological process or morphological characteristics that are not being examined as long as they do not appreciably affect the process or characteristic being examined.

5 **DESCRIPTION OF THE INVENTION**

This invention generally relates to novel compositions and methods for the treatment of certain cancers. Additionally, this invention relates to novel compositions and methods to screen drugs useful for the treatment of certain cancers.

10 Injury to DNA is minimized by enzymes that recognize errors, remove them, and replace the damaged DNA with corrected nucleotides. DNA damage occurs when a single-strand break is introduced, a base is removed leaving its former partner unpaired, a base is covalently modified, a base is converted into another that is not appropriately paired with the partner base, or a covalent link is introduced
15 between bases on opposite strands. Excision repair systems remove the mispaired or damaged base from the DNA strand and then synthesize new DNA to replace it. Base excision repair (BER) is initiated during replication of DNA and allows for correction of damaged bases/mispaired bases prior to completion of replication.

20 Base excision repair is initiated by a DNA glycosylase that removes N-glycosidic (base-sugar) bonds, liberating the damaged base and generating an abasic site (AP site). An apurinic or apyrimidinic site results from the loss of a purine or pyrimidine residue, respectively, from DNA. uracil residues result from the spontaneous deamination of cytosine and can lead to a C → T transition if unrepaired. There is also a glycosylase that recognizes and excises hypoxanthine,
25 the deamination product of adenine. Other glycosylases remove alkylated bases (such as 3-methyladenine, 3-methylguanine, and 7-methylguanine), ring-opened purines, oxidatively damaged bases, and in some organisms, UV photodimers.

30 The AP site is further processed by a 5'-3' endonuclease (AP endonuclease (APE)) that incises the phosphodiester bond on both sides of the damaged purine or pyrimidine base. The AP endonucleases introduce chain breaks by cleaving the phosphodiester bonds at the AP sites.

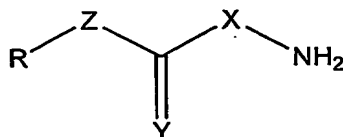
PARP aids in processing of DNA strand breaks induced during MMR and BER. PARP is a DNA nick surveillance protein that binds weakly to BER intermediates when single-nucleotide BER proceeds normally to completion. In contrast, when single nucleotide BER is stalled by a block in the excision step,
5 PARP binds strongly to the BER intermediate, along with AP endonuclease (APE), DNA pol β , and FEN-1.

In mammalian cells, the 5'-deoxyribose sugar phosphate is removed by the intrinsic AP lyase (dRP) activity of DNA polymerase β (pol β). DNA polymerase enzyme also fills the gaps with new nucleotides.

10 Finally, DNA ligase covalently links the 3' end of the new material to the old material. Thus, the wild-type sequence is restored.

Topoisomerases I and II are also involved in DNA repair, as they recognize spontaneous AP sites and form stable cleavable complexes. Topoisomerase II inhibitors promote DNA cleavage and other chromosomal aberrations, including
15 sister chromatid exchanges.

Thus, compounds useful as BER inhibitors include AP endonuclease inhibitors such as methoxyamine (MX), N-ethylmaleimide, O⁶-benzylguanine, and
20 compounds having structures of formula I:



Formula I

wherein X is O or NH,

Y is O, S, or NH,

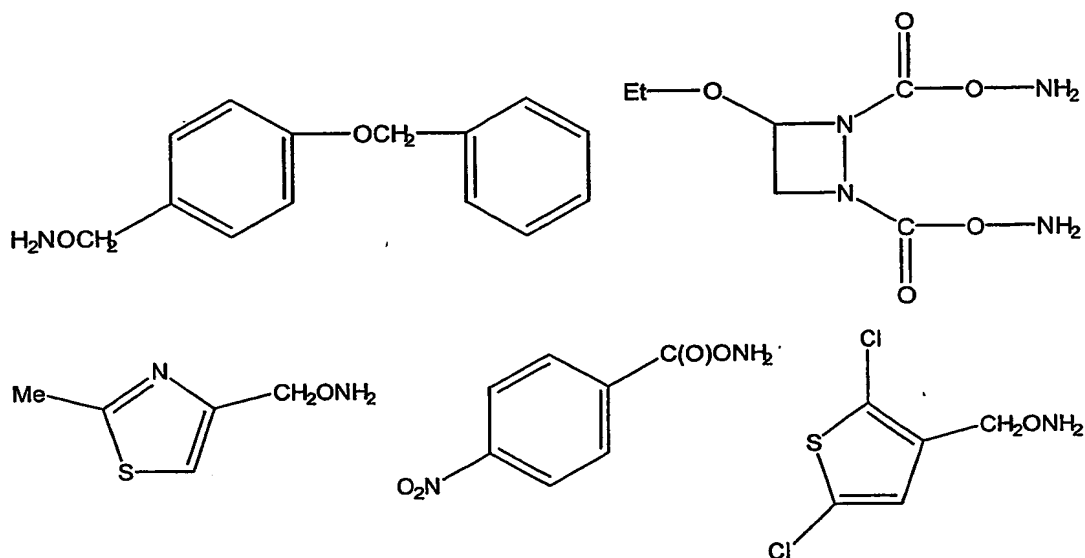
25 Z is absent or represents O, S, or NH, and
R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

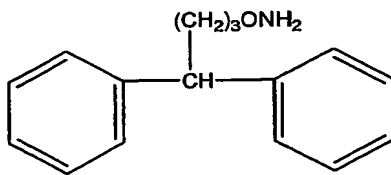
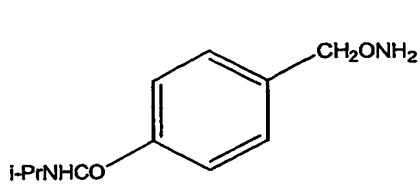
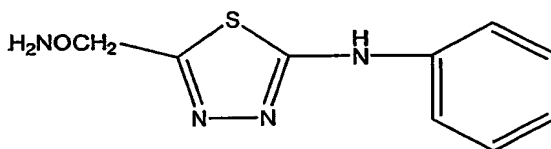
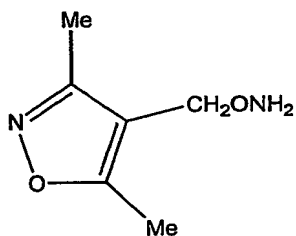
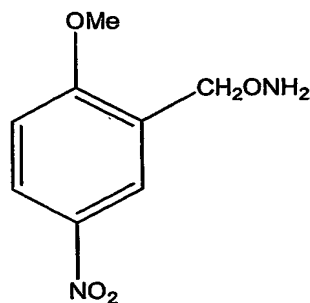
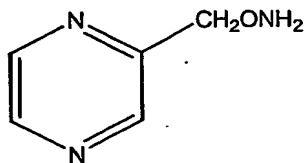
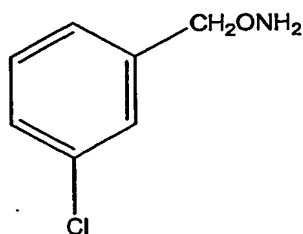
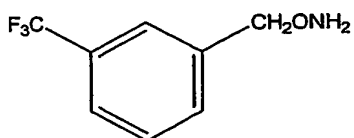
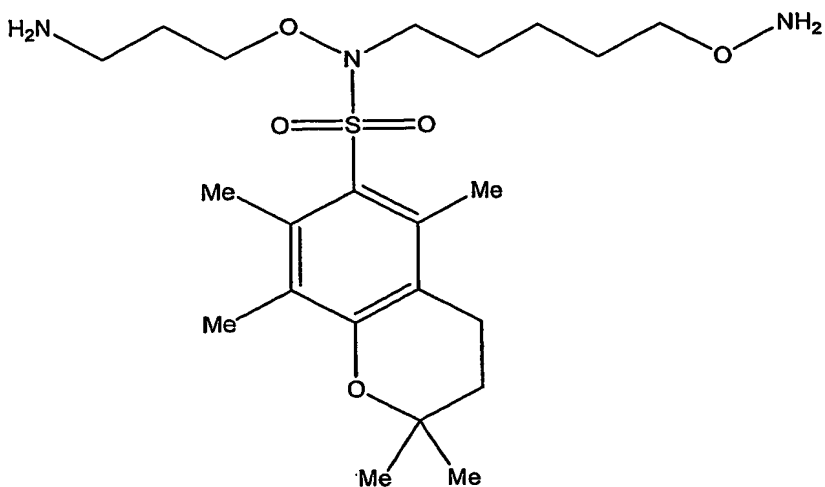
In single-nucleotide BER, the deoxyribose phosphate (dRP) in the abasic site is removed by the lyase activity of DNA pol β . Compounds such as methoxyamine
30 react with the aldehyde of an abasic site, making it refractory to the β -elimination

step of the dRP lyase mechanism, thus blocking single-nucleotide BER. Suitable AP endonuclease inhibitors may act by binding to AP sites and preventing APE-mediated cleavage of phosphodiester bonds, or by acting directly on AP endonuclease. Other compounds that may possess AP endonuclease inhibitory activity (e.g., by binding to AP sites and preventing APE-mediated cleavage of phosphodiester bonds) include Other potential inhibitors include O-benzylhydroxylamine; ethyl aminooxyacetate; aminooxyacetic acid; ethyl aminooxyacetate; $\text{H}_2\text{NOCHMeCO}_2\text{H}$; carboxymethoxyamine; aminooxyacetic acid; $\text{HN}=\text{C}(\text{NH}_2)\text{SCH}_2\text{CH}_2\text{ONH}_2$; $\text{H}_2\text{NO}(\text{CH}_2)_3\text{SC}(\text{NH}_2)=\text{NH}$;

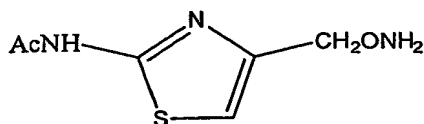
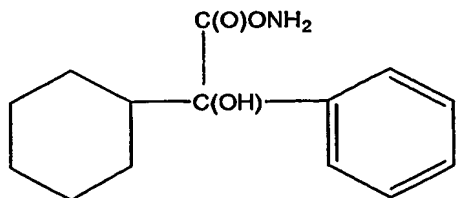
10 $\text{MeOC}(\text{O})\text{CH}(\text{NH}_2)\text{CH}_2\text{ONH}_2$; $\text{H}_2\text{NOCH}_2\text{CH}(\text{NH}_2)\text{CO}_2\text{H}$; canaline; $\text{H}_2\text{NO}(\text{CH}_2)_4\text{ONH}_2$; O-(p-nitrobenzyl)hydroxylamine; 2-amino-4-(aminooxymethyl)thiazole; 4-(aminooxymethyl)thiazole; O,O'-(o-phenylenedimethylene)dihydroxylamine; 2,4-dinitrophenoxyamine; O,O'-(m-phenylenedimethylene)dihydroxylamine; O,O'-(p-

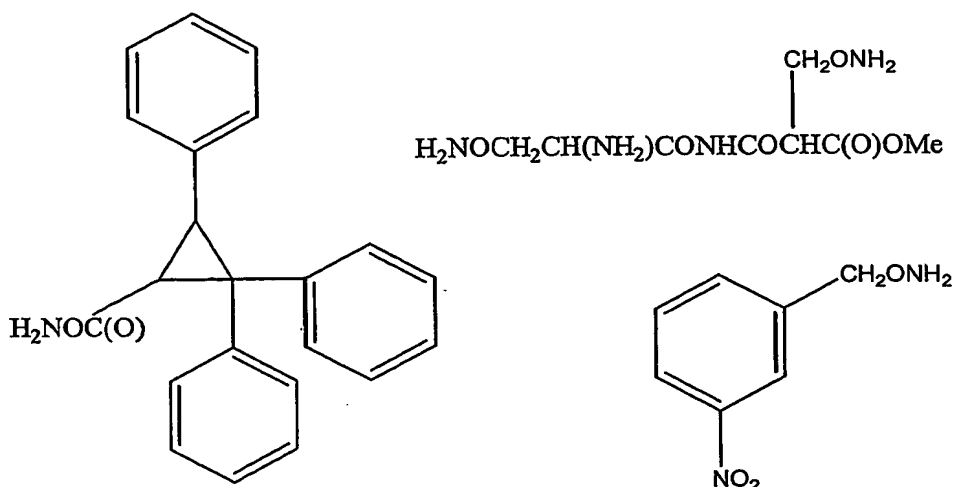
15 phenylenedimethylene)dihydroxylamine; $\text{H}_2\text{C}=\text{CHCH}_2\text{ONH}_2$; $\text{H}_2\text{NO}(\text{CH}_2)_4\text{ONH}_2$; $\text{H}_3\text{C}-(\text{CH}_2)_{15}-\text{O}-\text{NH}_2$, 2,2'-(1,2-ethanediyl)bis(3-aminooxy)butenedioic acid dimethyl diethyl ester; compounds having any of the following structures:





5





and pharmaceutically acceptable salts of any of these compounds.

Compounds useful as BER inhibitors include PARP inhibitors, such as 4-amino-1,8-naphthalimide (ANI), PD128763, 3-AB, 6-AN, and 8-hydroxy-2-methyl-quinazolin-4-[3H]one (NU-1025).

Compounds useful as BER inhibitors include DNA polymerase inhibitors (e.g., DNA polymerase β , γ , or ϵ), such as prunasin, aphidicolin, 2',3'-dideoxycytidine triphosphate (ddCTP), 2',3'-dideoxythymidine triphosphate (ddTTP), 2',3'-dideoxyadenosine triphosphate (ddATP), 2',3'-dideoxyguanosine triphosphate (ddGTP), 1-beta-D-arabinofuranosylcytosine (Ara-C), caffeine, arabinocytidine, and bleomycin.

Compounds useful as BER inhibitors include DNA ligase inhibitors (e.g., DNA ligase I, II, or III), such as ursolic and oleanolic acids, aleuritolic acid, protolichesterinic acid, swertifrancheside, fulvoplumierin, fagaronine chloride, and bleomycin. XRCC1 is the protein partner of DNA ligase III, and inhibitors of XRCC1, such as 3-AB, are useful as BER inhibitors as well.

Topoisomerase II inhibitors induce DNA cleavage and other chromosomal aberrations, including sister chromatid exchanges. Compounds useful as BER inhibitors also include topoisomerase II inhibitors, such as etoposide (VP-16, VP-16-123), meso-4,4'-(2,3-butanediyl)-bis-(2,6-piperazinedione) (ICRF-193, a bisdioxopiperazine), doxorubicin (DOX), amsacrine (4',9'-acridinylaminomethanesulfon-m-anisidide; mAMSA), pazelliptine, nalidixic acid, oxolinic acid, novobiocin, coumermycin A1, fostriecin, teniposide, mitoxantrone,

daunorubicin, N-[2-dimethylamino)ethyl]acridine-4-carboxamide (DACA), merbarone, quinacrine, ellipticines, epipodophyllotoxins, ethidium bromide, epirubicin, pirarubicin, 3'-deamino-3'-morpholino-13-deoxy-10-hydroxy carminomycin; 2'',3''-bis pentafluorophenoxyacetyl-4',6'-ethylidene-beta-D
5 glucoside of 4'-phosphate-4'-dimethylepipodophyllotoxin 2N-methyl glucamine salt (F11782; a fluorinated lipophilic epipodophyllotoxin), adriamycin, actinomycin D, anthracyclines (such as 9-aminoanthracycline), and pyrazoloacridine (PZA). Topoisomerase I inhibitors, such as camptothecin and topotecan can also be used as BER inhibitors.

10 Other inhibitors of these enzymes, whether known in the art or hereafter identified, as well as inhibitors of other elements of the BER pathway, such as DNA alkyltransferase, may be employed in the compositions and methods of the invention without departing from the scope and spirit of the present invention.

In certain embodiments, the present invention contemplates the use of 1) a
15 topoisomerase inhibitor, preferably a topoisomerase II inhibitor such as etoposide, 2) an anticancer agent that induces the formation of AP sites (such as temozolomide), and 3) a BER inhibitor (other than a topoisomerase inhibitor), such as methoxyamine.

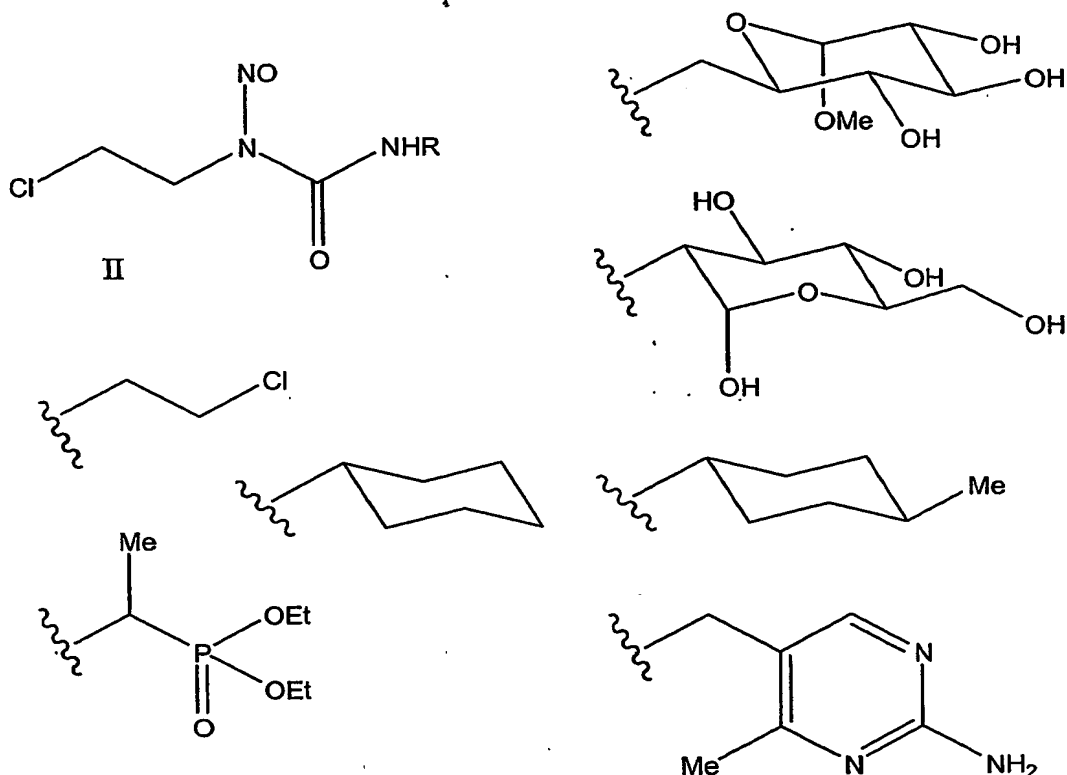
Anticancer agents that induce the formation of AP sites include intercalating
20 agents such as bleomycin, adriamycin, quinacrine, echinomycin (a quinoxaline antibiotic), and anthrapyrazoles.

Radiation, such as gamma radiation, UVA, and UVB, can be used to generate AP sites according to the methods of the invention. Ultraviolet light is absorbed in DNA with the formation of UV-specific di-pyrimidine photoproducts.
25 Exposure to gamma irradiation, UVA, and UVB can induce damaged pyrimidine photodimers

Anticancer agents that induce the formation of AP sites include DNA oxidizing agents such as hydrogen peroxide.

Anticancer agents that induce the formation of AP sites include alkylating
30 agents such as temozolomide (TMZ), 1,3-bis(2-chloroethyl)-1-nitrosourea (BCNU), MeOSO₂(CH₂)₂-leixitropsin (Me-Lex), cis-diamminedichloroplatinum II (cisplat; cis-DDP), mitomycin bioreductive alkylating agents, quinones, streptozotocin,

cyclophosphamide, nitrogen mustard family members such as chloroambucil, pentostatin (and related purine analogs), fludarabine, bendamustine hydrochloride, chloroethylating nitrosoureas (e.g., lomustine, fotemustine, cystemustine), dacarbazine (DTIC), and procarbazine. In certain embodiments, the alkylating agent
 5 is a nitrosourea such as a mustine, i.e., a compound having a structure of Formula II, wherein R is an optionally substituted hydrocarbon substituent, such as an alkyl, cycloalkyl, heterocyclyl, aryl, heteroaryl, cycloalkylalkyl, heterocyclalkyl, aralkyl, or a heteroaralkyl:



10

In preferred embodiments, R is a substituent shown below or to the right of Formula I, i.e., the chemotherapeutic is carmustine (BCNU), chlorozotocin, fotemustine, lomustine, nimustine, ranimustine, or semustine. In certain related embodiments, the chloroethyl group of Formula I is replaced by a methyl group, as in streptozocin. In
 15 certain embodiments, however, R is not 2-chloroethyl, i.e., the compound is not BCNU.

Alkylating agents can function by adding methyl groups to DNA, cross-linking macromolecules essential for cell division, and linking guanine bases in

DNA through their N^7 atoms. Both inter- and intra-strand cross-links can be mediated by alkylating agents. Inter-strand cross-links prevent the separation of the DNA strands necessary for cell division, and by being more difficult to repair, constitute the more lethal lesion.

5 In certain embodiments, the anticancer agent is selected from radiosensitizers such as 5-iodo-2'-deoxyuridine (IUdR), 5-fluorouracil (5-FU), 6-thioguanine, hypoxanthine, uracil, ecteinascidin-743, and camptothecin and analogs thereof.

In certain embodiments, the anticancer agent is not temozolomide. In certain embodiments, the anticancer agent is not BCNU. In certain embodiments, the
10 anticancer agent is not PE128723, 6-AN, 3-AB, BCNU, or temozolomide.

Throughout the specification, any recitation of a particular compound should be understood to encompass that compound and any (other) pharmaceutically acceptable salts thereof.

15 In DNA repair competent cells, DNA adducts formed by methylating agents may be efficiently repaired or may be sites of both mutagenic and cytotoxic damage. In this process, the cellular response is specific for each of the DNA adducts formed. Perhaps the best studied is the response to O^6 -methylguanine (O^6 mG). This adduct may be repaired in a single step reaction by O^6 -alkylguanine-DNA alkyltransferase
20 (AGT), however, saturation of this protein by an excess of adducts or inhibition by O^6 -benzylguanine (BG) results in residual adducts which are both cytotoxic and mutagenic (Pegg *et al.* "Structure, function and inhibition of O^6 -alkylguanine-DNA AGT" *Prog. Nucleic Acid Res. Mol. Biol.* 51:167-233, 1995.). Cytotoxicity results from recognition of this adduct by components of the mismatch repair (MMR)
25 system, a 5- or 6-protein complex which recognizes O^6 mG: thymine base pairs formed by DNA replication past O^6 mG, and excises thymine and surrounding bases, resulting in DNA strand breaks. However, a thymine is preferentially reincorporated opposite the persisting O^6 mG, triggering MMR function again. It has been
30 hypothesized that this repetitive aberrant repair process increases DNA double strand breaks and acts as a trigger of apoptosis (Fink *et al.* "The role of DNA mismatch repair in drug resistance" *Clin. Cancer Res.* 4:1-6, 1998.).

MMR deficiency results in inability to process the O⁶mG:T mispair. Consequently, cells replicate DNA past O⁶mG lesions without cell cycle arrest, chromosomal aberrations, or apoptosis and therefore survive in the face of persistent DNA damage (Branch *et al.* "Defective mismatch binding and a mutator phenotype in cells tolerant to DNA damage" *Nature* 362:652-654, 1993; Kat *et al.* "An alkylation-tolerant, mutator human cell line is deficient in strand-specific mismatch repair" *Proc. Natl. Acad. Sci., USA* 90:6424-6428, 1993; Karran and Bignami "DNA damage tolerance, mismatch repair and genome instability" *Bioessays* 16:833-839, 1994; Griffin *et al.* "DNA mismatch binding and incision at modified guanine bases by extracts of mammalian cells: implications for tolerance to DNA methylation damage" *Biochemistry* 33:4787-4793, 11994; Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide but not to 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56:5375-5379, 1996). The presence of MMR deficiency in a number of colon cancer cell lines allowed the present inventors the opportunity to evaluate the relative contribution of this DNA repair defect in resistance to the methylating chemotherapeutic agent, temozolomide (TMZ), where we found that MMR deficiency resulted in 35-60 fold resistance to TMZ in cells defective in either MLH1 or GTBP even after inhibition of AGT by BG (Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide by not to 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56:5375-5379, 1996).

While O⁶mG is the best studied cytotoxic DNA adduct, it is not the most abundant. TMZ, like other methylating agents, also forms N⁷-methylguanine (N⁷mG) and N³-methyladenine (N³mA) DNA adducts at frequencies 11 and 1.5 times that of O⁶mG. These DNA adducts are efficiently removed by base excision repair (BER) and appear to contribute little to cytotoxicity. In the first step of BER, a series of glycosylases recognize abnormal bases such as N³mA and N⁷mG (O'Connor *et al.* "Isolation and structure of a cDNA expressing a mammalian 3-methyladenine-DNA glycosylase" *EMBO J.* 9:3337-3342, 1990; Samson *et al.* "Cloning and characterization of a 3-methyladenine DNA glycosylase cDNA from human cells whose gene maps to chromosome 16" *Proc. Natl. Acad. Sci. USA* 88:9127-9131, 1991), the T:G mismatch (Neddermann *et al.* "Functional expression

of soluble human interleukin-11 (IL-11) receptor alpha and stoichiometry of in vitro IL-11 receptor complexes with gp130" *J. Biol. Chem.* 271:12767-12774, 1996), and deaminated bases such as hypoxanthine/oxidized 8-oxo-7,8-dihydroguanine or uracil:A (Vollberg *et al.* "Isolation and characterization of the human uracil DNA glycosylase gene" *Proc. Natl. Acad. Sci. USA* 86: 8693-8697, 1989; Olsen *et al.* "Molecular cloning of human uracil-DNA glycosylase, a highly conserved DNA repair enzyme" *EMBO J.* 8:3121-3125, 1989; Radicella *et al.* "Cloning and characterization of hOGG1, a human homolog of the OGG1 gene of *Saccharomyces cerevisiae*" *Proc. Natl. Acad. Sci. USA* 94:8010-8015, 1997; Rosenquist *et al.* "Cloning and characterization of a mammalian 8-oxoguanine DNA glycosylase" *Proc. Natl. Acad. Sci. USA* 94:7429-7434, 1997). Following enzymatic or spontaneous hydrolysis of the N-glycosidic bond and release of the abnormal base, AP (apurinic/apyrimidinic) endonuclease hydrolyzes the phosphodiester backbone 5' to the lesion and dRpase (a DNA deoxyribophosphodiesterase and its activity is associated with polymerase β) excises the residual dRp, generating a gap of one nucleotide. DNA polymerase β fills the gap and DNA ligase seals the nick. This pathway is called short-patch BER. An alternative pathway for BER involves DNA synthesis to fill a gap of 2 to 13 nucleotides. This long patch repair requires proliferating cell nuclear antigen (PCNA) and PCNA-dependent DNA polymerase (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407:203-215, 1998).

Poly-(ADP-ribose)-polymerase (PARP) acts as a nick sensor of DNA strand breaks by itself or interaction with XRCC1 and involves in BER. PARP binds damaged DNA, resulting in autoribosylation. The modified protein then releases and allows other proteins to access and repair DNA strand breaks (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407:203-215, 1998; Molinete *et al.* "Over production of the poly (ADP-ribose) polymerase DNA-binding domain blocks alkylation-induced DNA repair synthesis in mammalian cells" *EMBO J.* 12:2109-2117, 1993; Caldecott *et al.* "XRCCI polypeptide interacts with DNA polymerase β and possibly poly (ADP-ribose) polymerase, and DNA ligase III is a novel molecular 'nick-sensor' in vitro" *Nucleic Acids Res.* 24:4387-4394, 1996). Therefore, PARP participates in BER after nick formation in both short

patch and long patch repair. It appears most active in the alternative (long patch repair) pathway for BER.

BER as a therapeutic target to increase the cytotoxicity of methylating agents has been documented. Cells deficient in DNA polymerase B or blocked in
5 expression of AP endonuclease by antisense oligonucleotides are sensitized to methylating agents (Sobol *et al.* "Requirement of mammalian DNA polymerase- β in base-excision repair" *Nature* 379:183-186, 1996; Walker *et al.* "A role for the human DNA repair enzyme HAP1 in cellular protection against DNA damaging agents and hypoxic stress" *Nucleic Acids Res* 22:4884-4889, 1994). In addition,
10 mice deficient in N³-methyladenine DNA glycosylase exhibited increased sensitivity to alkylating drugs like BCNU and mitomycin C (Engelward *et al.* "Repair-deficient 3-methyladenine DNA glycosylase homozygous mutant mouse cells have increased sensitivity to alkylation-induced chromosome damage and cell killing" *EMBO* 15:945-952, 1996). On the other hand, overexpression of the N³-methyladenine
15 DNA glycosylase, which increases the number of AP sites formed, also increases the cytotoxicity of methylating agents (Coquerelle *et al.* "Overexpression of N-methylpurine-DNA glycosylase in Chinese hamster ovary cells renders them more sensitive to the production of chromosomal aberrations by methylating agents - a case of imbalanced DNA repair" *Mutation Res.* 336:9-17; 1995.). Finally, cells
20 lacking PARP activity are more sensitive to alkylating agents, with increased apoptosis and chromosomal instability (Menissier-de Murcia *et al.* "Requirement of poly (ADP-ribose) polymerase in recovery from DNA damage in mice and in cells" *Proc. Natl. Acad. Sci. USA* 94:7303-7304, 1997; Boulton *et al.* "Potentiation of temozolomide-induced cytotoxicity: a comparative study of the biological effects of poly (ADP-ribose) polymerase inhibitors" *Br. J. Cancer*, 72:849-856, 1995). These
25 data suggest that balanced expression of proteins in the base excision repair complex is important to the efficient processing of lesions. BER is an important mechanism of resistance to therapeutic methylating agents. Accordingly, the present invention contemplates compositions and methods that enhance the effect of a DNA-damaging
30 chemotherapeutic agent (e.g., an agent that induces the formation of AP sites in a cell) through the use of a BER inhibitor in conjunction with the anticancer agent.

We examined two classes of agents which could inhibit the BER pathway to determine whether they would increase the cytotoxicity of methylating agents in colon cancer cells, particularly in cells deficient in MMR. Since MMR-deficient cells are tolerant to O⁶mG formed by TMZ, any change in cytotoxicity observed after use of a BER inhibitor would be due to interruption in repair of N⁷mG and N³mA DNA adducts. Our first strategy was to combine methoxyamine (MX) with TMZ. MX has been shown to react with the free aldehyde formed at the abasic site exposed by glycosylases and to reduce cleavage at AP sites in mammalian cells, suggesting that the MX-bound abasic site is not a substrate for AP endonuclease (Fortini *et al.* "Mutagenic processing of ethylation damage in mammalian cells: the use of methoxyamine to study apurinic/apyrimidinic site-induced mutagenesis" *Cancer Res.* 53:1149-1155, 1993). Since AP sites modified by MX are relatively stable and must be converted to cytotoxic lesions, we hypothesized that MX would interrupt BER in cells and potentiate the cytotoxic effects of TMZ, even in MMR defective cells. The second strategy we used was to inhibit PARP with 3,4-dihydro-5-methoxyisoquinolin-1(2H)-one (PD128763), 3-aminobenzamide (3-AB) or 6-aminonicotinamide (6-AN) and to subsequently treat cells with TMZ. We hypothesized that inactivated PARP would affect short- and long-patch BER, destabilize strand breaks, reduce interaction with other proteins during repair of methylated DNA adducts, and lead to cell death in both MMR-proficient and -deficient cells.

As contemplated above, since MMR-defective cell lines are remarkably resistant to methylating agents yet accumulate high levels of 3 methylating DNA adducts, O⁶mG, N³mA and N⁷mG, we reasoned that the interruption of repair of N³mA and N⁷mG adducts by the BER process would sensitize cells to methylating agents. To address this issue, we studied the effect of MX on potentiation of TMZ-induced cytotoxicity. MX interacts specifically with the tautomeric open-ring form of deoxyribose generated from the removal of an abnormal base by glycosylase. The MX-modified AP site is relatively stable (Liuzzi and Talpeart-Borle "A new approach to the study of the base-excision repair pathway using methoxyamine" *J. Biol. Chem.* 260:5252-5258, 1985; Rosa *et al.* "Processing in vitro of an abasic site reacted with methoxyamine: a new assay for the detection of abasic sites formed in

vivo" *Nucleic Acids Res.* 19: 5569-5574, 1991) and inhibits the cleavage of AP sites in DNA by AP endonuclease in mammalian cells. This has been shown to protect cells from cytotoxicity and mutagenicity induced by SN₁-type ethylating agents, such as ENU, but not SN₂ alkylating agents diethylsulfate and MMS (Fortini *et al.* "Mutagenic processing of ethylation damage in mammalian cells: the use of methoxyamine to study apurinic/apyrimidinic site-induced mutagenesis" *Cancer Res.* 53: 1149-1155, 1993; Fortini *et al.* "Methoxyamine modification of abasic sites protects CHO cells from the cytotoxic and mutagenic effects of oxygen alkylation" *Carcinogenesis* 13: 87-93, 1992). Moreover, the protection was strictly time-dependent and was limited to the short period (30 min) after exposure to the alkylating agents (Fortini *et al.* "Methoxyamine modification of abasic sites protects CHO cells from the cytotoxic and mutagenic effects of oxygen alkylation" *Carcinogenesis* 13: 87-93, 1992).

In our initial studies, we observed that MX reduced cleavage at AP sites and decreased BER in human colon cancer cell extracts. However, we did not see protection of these two cell lines from ENU cytotoxicity using longer duration exposure to MX. The short duration of MX studied previously may not have the same impact on BER inhibition as does a longer exposure to MX. Our results showed that MX synergistically increased TMZ-induced cytotoxicity in human colon cancer cell lines in both MMR-proficient and -deficient cells. A similar degree of enhanced cytotoxicity was observed with MX and MMS and with TMZ as well. The effect of BG inhibition of AGT was additive to the effect of MX only in the MMR-proficient SW480 cell line but not in the MMR defective HCT116 cell line. These data suggest that O⁶mG DNA adducts do not contribute to the enhanced cytotoxic effect of TMZ by MX. Furthermore, a similar degree of enhanced cytotoxicity was observed with MX and MMS as with TMZ, again implicating N³nA and N⁷mG induced abasic sites as the major targets for MX. In our recent studies, a prolonged exposure to low dose MX results in even greater potentiation of TMZ cytotoxicity.

Without wishing to be bound by any theory, mechanisms of MX-enhanced cytotoxicity of methylating agents in colon cancer cells may be due to i) the MX-AP site complex being able to block the AP endonucleolytic step of the BER pathway;

ii) the persistence of abasic sites increasing topoisomerase II-mediated DNA cleavage (Kingma and Osheroff "Apurinic sites are position-specific topoisomerase II poisons" *J. Biol. Chem.* 272: 1148-1155, 1997); iii) AP sites inhibiting DNA replication and triggering programmed cell death (Robertson *et al.* "Down-
5 regulation of apurinic/aprimidinic endonuclease expression is associated with the induction of apoptosis in differentiating myeloid leukemia cells" *Cell Growth & Differentiation* 8: 443 -449, 1997).

Under normal circumstances, TMZ produces strand breaks during BER-mediated repair of N⁷mG and N³mA adducts which are efficiently repaired and do
10 not contribute to cytotoxicity until high concentrations of adducts are achieved. When DNA strand breaks are present, one component of the response is recognition, binding and activation of PARP. Activated PARP leads to autoribosylation, and this in turn facilitates access of repair enzymes to DNA damage (Willson "Mammalian base excision repair and DNA polymerase beta" *Mutation Res.* 407: 203-215, 1998;
15 Buschfort *et al.* "DNA excision repair profiles of normal and leukemic human lymphocytes: functional analysis at the single-cell level" *Cancer Res.* 57: 651-658, 1997) and appears to enhance processing of stand breaks and religation by polymerase-β and ligase I (Althaus *et al.* "Histone shuttle driven by the automodification cycle of poly(ADP-ribose) polymerase" *Environ Mol Mutagen*, 22:
20 278-282, 1993). In the alternative BER pathway, PARP interacts with XRCC1 to facilitate repair (Caldecott *et al.* "XRCC 1 polypeptide interacts with DNA polymerase β and possibly poly (ADP-ribose) polymerase, and DNA ligase III is a novel molecular 'nick-sensor' in vitro" *Nucleic Acids Res.* 24: 4387-4394, 1996). It seems likely that PARP plays an important role in communication between repair
25 proteins and stability of the repair complex involved in BER. This suggests that inhibition of PARP leads to an impaired ability to rejoin DNA strand breaks which can initiate both apoptotic and non-apoptotic cell death cascades and thereby increase cytotoxicity of TMZ (Wedge *et al.* "3-Aminobenzamide and/or O⁶benzylguanine evaluated as an adjuvant to temozolomide or BCNU treatment in
30 cell lines of variable mismatch repair status and O⁶-alkylguanine-DNA alkyltransferase activity" *Br. J. Cancer* 74: 1030-1036, 1996). Our results support

this hypothesis. Potentiation of cytotoxicity of methylating agent with PARP inhibitors was observed with a marked increase in apoptosis and PARP cleavage.

Results of BCNU combined with either MX or PD128763 are in sharp contrast to TMZ; little if any potentiation is observed in the absence of BG in either cell line. This suggests that while BER appears to process BCNU induced crosslinks, inhibition of BER in this manner has little impact on BCNU toxicity. One of the best studied BCNU-induced lethal lesions is the N³C-N¹G interstrand cross-link formed after initial chloroethyl monoadducts at O⁶G and cyclic rearrangement to N¹,O⁶-ethanoguanine (Wilson *et al.* "Life without DNA repair" *Proc. Natl. Acad. Sci. USA* 94: 7303-7304, 1997). However, treatment of cells with BCNU also produces alkylated bases that may be labile and spontaneously result in breakage or nicking of the phosphoribosyl backbone (Gonzaga *et al.* "Identification of the cross-link between human O⁶-metnine-DNA methyltransferase and chloroethylnitrosourea-treated DNA" *Cancer Res.* 52:6052 -6058, 1992). Since PARP has been shown to bind to BCNU-induced DNA nicks *in vitro* (Prakash and Gibson "Sequence-selective depurination, DNA interstrand cross-linking and DNA strand break formation associated with alkylated DNA" *Carcinogenesis* 13: 425-431, 1992), it is reasonable to assume that PD128763 might increase BCNU-mediated cytotoxicity. However, our studies showed only minor enhancement of toxicity in HCT116 cells and no enhancement in SW480 cells. Even though methyladenine DNA glycosylase has been implicated in BCNU cross-link repair and its absence sensitizes cells to BCNU (Malapetsa *et al.* "Identification of a 116 kDa protein able to bind 1,3-bis (2-chloroethyl)-1-nitrosourea-damaged DNA as poly (ADP-ribose) polymerase" *Mutation Res.* 362: 41-50, 1996), we did not observe sensitization to BCNU by treatment with MX in the absence of BG. In the presence of BG, MX potentiated BCNU toxicity, indicating that MX may interfere with DNA cross-link repair pathway and suggesting that BER may be involved in repair of the N¹O⁶anoguanine cross-link which is not formed if AGT reacts with the O⁶chloroethylguanine adduct. Taken together, these data suggest a different reaction of MX with damaged DNA induced by BCNU compared to TMZ. With TMZ, MX-enhanced cytotoxicity is associated with AP sites generated from repair of N⁷mG

and N³mA DNA adducts formed by methylating agent but with BCNU, it might be the O⁶ lesion-induced cross-link which controls BCNU toxicity.

While an understanding of the precise mechanism is not needed to practice the present invention, it appears that apoptosis mediates both MX and PD128763-enhanced cytotoxicity of TMZ. Increased apoptosis was observed in MMR wt
5 SW480 cells but not in MMR-deficient HCT116 cells after treatment with BG and TMZ. This suggests that MMR processing of O⁶mG is a potent apoptosis-inducing event (Matijasevic *et al.* "Protection against chloroethylnitrosourea cytotoxicity by eukaryotic 3-methyladenine DNA glycosylase" *Proc. Natl. Acad. Sci. USA*,
10 90:11855-11859, 1993). Although the biological and functional consequences of PARP and its cleavage in apoptosis still remain to be further identified, it has been demonstrated that PARP is rapidly and specifically cleaved during apoptosis (Kaina *et al.* "Chromosomal instability, reproductive cell death and apoptosis induced by O⁶-methylguanine in Mex-, Mex+ and methylation-tolerant mismatch repair
15 compromised cells: facts and models" *Mutation Res.* 381: 227-241, 1997; Lindahl *et al.* "Post-translational modification of poly(ADP-ribose) polymerase induced by DNA strand breaks" *Trends Biochem Sci* 20: 405-411, 1995). PARP cleavage was observed in both SW480 and HCT116 cells after treatment with either MX or one of the PARP inhibitors and TMZ, confirming activation of apoptotic pathways.

20 We noted that arrest at cell cycle checkpoints paralleled the cellular response to DNA damage and that these were dependent on MMR and BER pathways. MMR wt SW480 cells were sensitive to TMZ alone with arrest in the S and G2 phases (Fink *et al.* "The role of DNA mismatch repair in drug resistance" *Clin. Cancer Res.* 4: 1-6, 1998). The S and G2 arrest were potentiated by MX or by PD128763 despite
25 the fact that SW480 is a p53 mutant cell line. In contrast, even high levels of DNA adducts formed by TMZ in the MMR deficient HCT116 cells did not induce cell cycle checkpoint arrest despite the fact that p53 is wild type in this cell line. This dysregulation of damage-induced cell cycle checkpoint control appeared due to failure of processing O⁶mG lesions in MMR deficient cells. However after combined
30 treatment with TMZ and either MX or PD 128763, HCT 116 cells showed S/G2 phase arrest and apoptosis. These results are consistent with previous studies of cell cycle changes following MMS exposure or other compounds that produce 90%

N³mA (Lazebnik *et al.* "Cleavage of poly(ADP-ribose) polymerase by a proteinase with properties like ICE" *Nature* 371: 346-347, 1994; Morris *et al.* "Flow cytometric evaluation of cell-cycle progression in ethyl methanesulfonate and methyl methanesulfonate-exposed P3 cells: relationship to the induction of sister-chromatid exchanges and cellular toxicity" *Environ. Mol. Mutagen.* 18: 139-149, 1991) and the prolonged G2 arrest observed in PARP knockout mice or derived cell lines (Menissier-de Murcia *et al.* "Requirement of poly (ADP - ribose) polymerase in recovery from DNA damage in mice and in cells" *Proc. Nat. Acad. Sci. USA* 94: 7303-7304, 1997) following DNA damage. These data indicate that both SW480 and HCT116 cells have a similar response to persistent N⁷mG and N³mA lesions, following interruption of BER.

Again, while an understanding of the mechanism is not necessary to practice the invention, it appears that disrupted BER processing of non-O⁶mG, most likely N⁷mG and N³mA DNA adducts formed by TMZ is cytotoxic to colon cancer cell lines. This may be particularly important in MMR deficient cells which are resistant to TMZ alone due to the failure to recognize O⁶mG DNA adducts. These studies provide evidence that disrupting repair of N⁷mG and N³mA by inhibiting BER or PARP may improve the therapeutic efficacy of methylating agents.

We also investigated whether TMZ and MX-induced MX-AP sites may also act as poisons of topoisomerase II α (topo II), producing dual lethal targets. To analyze the effect of MX-AP site on topo II-mediated DNA cleavage, a position-specific MX-AP site was generated on topo II cleavage sites using oligonucleotides for cleavage assay. Results showed that MX-AP sites were cleaved by both topo II and nuclear extraction from tumor cells, suggesting that MX-AP sites, when located within a topo II cleavage site, stimulate topo II-mediated cleavage, which has been referred to the action of topo II poison. We also found prolonged induction of topo II and remarkable increase in DNA double strand breaks in cells treated with TMZ and MX compared to TMZ alone. Similar results were also observed in cells treated with etoposide but not with N-methyl-maleimide (a catalytic inhibitor of topo II and also an antagonist of etoposide). Moreover, the potentiation of TMZ by MX was decreased in DEtp cell line that was a subline derived from DLD1 and 5-fold more resistant to etoposide compared to the parental cell line. These findings demonstrate

that treatments with TMZ combined with MX exert the same effect as etoposide does to trap or stabilize topo II in DNA cleavage complex that leads to the accumulation of cytotoxic DNA breaks.

5 Generally, the nomenclature used hereafter and the laboratory procedures in cell culture, tissue culture, tumor biology, and molecular genetics described below are those well known and commonly employed in the art. Standard techniques are used for cell culture methods, experimental design and compound formulation and nomenclature. Generally chemical reactions and purification steps are performed
10 according to the manufacturer's specifications. The techniques and procedures are generally performed according to conventional methods in the art and various general references (see, generally, Sambrook *et al.* *Molecular Cloning: A Laboratory Manual*, 2d ed. (1989) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., and *Current Protocols in Molecular Biology* (1996) John Wiley and Sons, Inc.,
15 N.Y., which are incorporated herein by reference) which are provided throughout this document. All the information contained therein is incorporated herein by reference. Assays for detecting the ability of agents to inhibit or enhance TMZ-mediated tumor reduction provide for facile high-throughput screening of agent banks (e.g., compound libraries, peptide libraries, and the like) to identify
20 potentiators. Such TMZ potentiators may be further developed as potential therapeutics and/or diagnostic or prognostic tools for diverse types of cancers. Likewise, TMZ and various potentiators may be useful in the treatment or diagnostics of other diseases. For example, such combinations of drugs may be directed to virally or parasitically infected cells.

25 Thus, the present invention provides a method for potentiating the toxicity of temozolomide in human cancer cells by administering an APE- or a PARP-inhibiting amount of an APE or PARP inhibitor. The present invention also provides a product comprising temozolomide and an APE or PARP inhibitor as a combined preparation for simultaneous, separate or sequential administration in said treatment
30 of human cancer cells. Additionally, the present invention provides a method for screening compounds that may potentiate the toxicity of temozolomide by inhibiting APE or PARP, or by other mechanisms.

I. Potentiation of Temozolomide Treatment for Cancer

The present invention contemplates compositions and methods for the potentiation of the elimination of cancer cells by temozolomide. While not limiting the invention to any particular mechanism, it is believed that temozolomide works as an anticancer agent by producing the DNA adducts O⁶mG, N⁷mG and N³mA. An accumulation of adduct results in the apoptotic elimination of the cell. However, the O⁶mG adducts are typically repaired by the MMR system and the N⁷mG and N³mA adducts are typically repaired by the base excision repair (BER) system. After removal of the abnormal base by methylpurine glycosylase, AP endonuclease (APE) cleaves the back bone and removes the sugar, allowing DNA repair. If the number of adducts produced outstrips the cells ability to repair the damage, or the BER repair mechanism is rendered ineffective, damage will accumulate and the cell will be eliminated through an apoptotic mechanism. Methoxyamine reacts with the abasic site and prevents APE cleavage, disrupting DNA repair. Methoxyamine has been shown in the present invention to potentiate the activity of temozolomide in the treatment of cancer. PARP aids in the repair of DNA strand breaks induced during MMR and BER. As such, inhibitors of PARP activity have been shown in the present invention to be effective in potentiating the activity of temozolomide in the elimination of cancer cells. The effectiveness of temozolomide as an anticancer agent can be greatly enhanced when used in conjunction with a potentiator that prevents the repair of the DNA adducts created as a result of treatment of the cell with temozolomide.

Temozolomide has been found to be most preferably administered in repeat dosages on consecutive days, and the dramatic potentiation effects of the present invention are realized in the highly preferred regimen involving the administration of MX or a PARP inhibitor prior to, or concurrent with, the administration of each dose.

Preferably, this administration of an APE or PARP inhibitor is repeated over a period of several days and is prior to the administered doses of temozolomide. Repeat doses can be administered at 1, 2, 3, 4 or 5 days or more with the preferred period of therapy determined by the response of the tumor to the treatment.

In a preferred embodiment, the APE or PARP inhibitor is administered in an APE- or PARP-inhibiting amount, *i.e.*, an amount sufficient to sensitize the tumor *in vivo* without causing undue sensitization of normal tissue, when the APE or PARP inhibitor is used concurrently with temozolomide.

5 The amount of the APE or PARP inhibitor used in the present invention varies according to the degree of the effective amount required for the treating of tumor cells. A suitable dosage is that which will result in a concentration of the APE or PARP inhibitor in the tumor cells to be treated which results in the depletion of the APE or PARP activity, respectively. If desired, some tumor cells can be obtained
10 by biopsy and can be tested *in vitro* for their sensitivity prior to *in vivo* treatment.

The neoplasms for which temozolomide is a particularly suitable treatment include carcinomas, melanomas, sarcomas, lymphomas and leukemias, with specific utility for astrocytoma, gliomas, malignant melanoma, chronic lymphocytic leukemia, lung cancer and best(sic) cancer.

15 Typical dosage ranges of temozolomide are generally between 0.1 and 200, preferably between 1 and 20 mg/kg body weight per day. The amount of APE inhibitor or PARP inhibitor necessary for the potentiation of temozolomide is dependent upon the amount of APE or PARP, respectively, normally present in the cancer cell type. A cancer cell having higher levels of APE or PARP may be
20 affected more dramatically by the preadministration or simultaneous administration of the APE or PARP inhibitor. Additionally, the amount of APE or PARP inhibitor necessary depends on the age and condition of the patient, the severity of the cancer being treated and the particular inhibitor being utilized.

II. Drug Screens

25 As noted above, the present invention contemplates screening assays for identifying compounds that inhibit tumor growth. The present invention contemplates a screening assay utilizing cell lines that are resistant to an anticancer drug such as temozolomide to screen for compounds that potentiate the anticancer drug. In one embodiment, a resistant cell line is pretreated with the test compound
30 followed by treatment with the anticancer agent. Cell growth and/or apoptosis are then measured. In another embodiment, the test compound is given simultaneously with the anticancer agent.

Additionally, the present invention contemplates the screening of compounds by the use of xenographs. In one embodiment, tumor cells are injected into immune-compromised mice (or other suitable animal) and allowed to grow. Mice carrying the tumors are then treated with the test compound in parallel with or prior to
5 treatment with an anticancer agent such as temozolomide. Tumor size is then measured to follow the effect of the treatment.

It is not intended that the present invention be limited by the nature of the drugs screened in the screening assay of the present invention. A variety of compounds, including peptides, organic compounds, nonorganic compounds, as well
10 as, formulations of more than one compound, are contemplated.

It is also not intended that the present invention be limited by the particular tumor cells used for the drug testing. A variety of tumor cells (for both positive and negative controls) are contemplated (including but not limited to cells set forth
above and in the examples below).

15 It is also not intended that the present invention be limited by the mechanism by which the potentiation is induced. For example, the compound may work by inhibiting APE cleavage, by inhibiting other DNA repair mechanisms (*e.g.*, PARP) or by other as of yet unidentified mechanisms.

It is contemplated that the invention be utilized in the screening of
20 compounds by high-throughput screening methods. For example, automated systems and microscale assay systems are contemplated. Furthermore, it is contemplated that the invention can be used for the screening of compound libraries (*e.g.*, drug libraries) utilizing these high-throughput screening methods.

The identification of BER inhibitors as enhancers of anticancer drugs is
25 important because it identifies a new target, AP sites in DNA for anticancer drug development. Recent evidence indicates that AP sites are common DNA lesions for anticancer drugs that attack DNA, generating modified bases that are removed by DNA glycosylases. Thus, BER inhibitors have the potential to improve the therapeutic efficacy of a broad spectrum of anticancer agents. MX is an effective
30 prototypical agent and leads to the direction for the development of new drugs. Below is a high-throughput screening assay to identify new inhibitors of BER that

are able to block AP site cleavage by APE and potentially enhance therapeutic efficacy. Screening studies will provide a broad effort at new drug development.

High-throughput screening methods include two molecular reaction assays:

1. Analysis of chemical - modified AP Sites assayed by Aldehyde Reactive Probe (ARP). This is a competitive assay to measure the reactivity with AP site
5 between ARP reagent (Dojindo Molecular Technologies Inc., Gaithersburg, MD) and the screening compounds. ARP and MX have a similar reactivity with AP sites. They react specifically with an aldehyde group that is open ring form of the AP sites. Thus this assay will allow identification of compounds with potential to block
10 AP site repair based on the binding affinity and efficiency to AP sites of screening compounds compared to ARP and MX.

a. AP site standard preparation: AP sites were produced in a calf thymus DNA by heat/acid-buffer solution. Intact calf thymus DNA was added to sodium citrate buffer (10 mM sodium citrate containing 10 mM NaH_2PO_4 and 10 mM NaCl,
15 pH 5.0) and held at 70 °C for 30 min. The reaction was stopped by chilling rapidly on ice, and the DNA was then precipitated with cold ethanol, washed with 70% ethanol, dried, and resuspended in sterilized distilled water.

b. AP-DNA (15 pg) was incubated with test compounds at different concentrations at 37 °C for 30 min prior to ARP (1 mM) or ARP alone (Dojindo
20 Molecular Technologies Inc., Gaithersburg, MD) for 30 min. After precipitation and wash with ethanol, DNA was resuspended in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH 7.2). DNA was heat-denatured at 100 °C for 5 min, quickly chilled on ice, and mixed with an equal amount of 2 M ammonium acetate. The single-stranded DNA was then immobilized on a BAS-85 NC membrane (Schleicher and Schuell)
25 using a vacuum filter device (Schleicher and Schuell). The NC membrane was incubated with streptavidinconjugated horseradish peroxidase (BioGenix) at room temperature for 30 min. After NC membrane was rinsed with washing buffer containing NaCl (0.26 M), EDTA (1 mM), Tris-HCl (20 mM), and Tween 20 (1%), ARP-AP sites are visualized with ECL reagents (Amersham Corp.) (Figure 22A)
30 and quantitated by scanning densitometer (Figure 22B).

2. AP sites cleaved by AP-endonuclease (APE). This assay confirms that AP sites modified by potential BER inhibitors are resistant to cleavage by APE,

(Trevigen, Gaithersburg, MD) a BER protein. The assay may be performed as follows (see also Figure 23A and B):

- a. AP site is prepared by replacing single nucleoside with deoxyuridine in duplex oligonucleotides (40 mer).
- 5 b. Regular AP site is produced in the duplex oligonucleotides by human uracil DNA glycosylase (LIDGase, Trevigen, Gaithersburg, MD) to remove the uracil residue.
- c. To generate MX-adducted AP site substrates: the UDG-treated duplex oligonucleotides are mixed with 10 mM MX in buffer containing 10 mM
10 KPO₄, pH 7.1 and incubated at 37 °C. After 30 min, the substrates are recovered by ethanol precipitation, lyophilized, resuspended in water, and stored at -20 °C.
- d. APE-cleavage reaction: DNA substrates containing either regular AP-sites or chemicalmodified AP sites are incubated with APE (Trevigen, Gaithersburg, MD) for 30 min and reactants are precipitated with 100% cold ethanol, washed with
15 70% ethanol and resuspended in TE buffer. The reactants are resolved by denaturing 20% polyacrylamide gel electrophoresis and visualized by silver staining (Silver Staining Kit, Pharmacia Biotech).

EXEMPLIFICATION

20

Materials and Methods

Chemicals and Reagents. BG was generously provided by Dr. Robert Moschel (Frederick Cancer Research and Development Center, National Cancer Institute). Stock solution was made in dimethylsulfoxide. TMZ and BCNU were obtained from the Drug Synthesis and Chemistry Branch, Drug Therapeutic Program, National
25 Cancer Center Institute. PD128763 was a gift from Park-Davis Pharmaceutical Division. 6-AN, 3-AB, MX and methylmethane sulfonate (MMS) were purchased from Sigma (ST. Louis, MO). Stock solutions of PD128763, 3-AB and 6-AN were prepared by dissolving in dimethylsulfoxide and added to cell culture at a final concentration of <1% DMSO, when cells were treated with these compounds. MX
30 was dissolved in sterilized water (pH 7.0). All stock solutions were kept at -20 °C. BCNU was prepared fresh in 0.5 ml of 100% ethanol, diluted in PBS, and used with in 10 min.

Colony Survival Assay. SW480 cells were obtained from the ATCC. HCT116 cells were obtained from R. Boland, University of Michigan Medical Center. All cell lines were cultured in appropriate growth medium.

5 Cells (2000/dish) were plated, adhered for 18 hrs and treated with TMZ or
 MMS plus or minus modifiers such as BX, MX, 6-AN, 3-AB or PD128763,
 according to experimental protocol. After treatment, cells were washed and fresh
 medium was replaced. The cells were grown for a further 7 days prior to staining
 with methylene blue for determination of colonies containing more than 50 cells.
10 Comparisons of drug induced cytotoxicity consisted of a calculation of the dose
 modification factor (DMF), defined as the ratio of the IC_{50} of either TMZ or MMS
 in the absence to that in the presence of indicted modifier(s), i.e., $DMF = IC_{50}$ for
 TMZ alone/ IC_{50} for TMZ plus modifier(s).

15 **Median Effect Analysis (MEA).** MEA was used to determine the dose-response
 interactions between TMZ and either MX or PD 128763. Drugs were combined at
 the ratio of the IC_{50} values for either TMZ and MX or TMZ and PD 128763 as
 determined by survival / concentration curves. The combination was compared to
 the cytotoxicity of each drug alone in every experiment. The combination index (CI)
20 was determined from colony-forming assays at increasing levels of cell killing,
 using analysis of multiple drug interaction program (Biosoft, Cambridge, United
 Kingdom) developed based on method of Chou and Talalay (Chou and Talalay
 "Quantitative analysis of dose-effect relationship: the combined effects of multiple
 drugs on enzyme inhibitors" In: G. Weber (ed.) *Advances in Enzyme Regulation*, pp.
25 27-55, New York: Pergamon Press, 1983). CI values of less than or grater than 1
 indicate synergy and antagonism, respectively., whereas a CI value of 1 indicates
 additivity of the drugs.

30 **Flow Cytometry for Cell Cycle Distribution Analysis.** For cell cycle analysis, 10^6
 cells were plated in 100-mm tissue culture dishes and exposed to MX (6
 mM)/PD128763 (100 μ M) or MX (6 mM)/PD128763 (100 μ M) plus TMZ (300
 μ M) at 37 °C. After 24 h-72 h of culture, cells were fixed in 80% ethanol and DNA

was stained with 20 µg/ml propidium iodide. DNA fluorescence of PI-stained cells was measured with an Elite ESP flow cytometer/cell sorter (Coulter, Miami, FL). Cell cycle distribution was analyzed with Modfit 5.2 program (Verity software, Topsham, MA) at least 10,000 cells per data point.

5

Western Blotting for PARP Cleavage Detection. Cell extracts were resolved by SDS-PAGE (12% polyacrylamide) in a Bio-Rad minigel apparatus at 150 V for 1 hr. Proteins were transferred onto PVDV membranes, using a Bio-Rad mini Trans-Blot cell for 1 hr at 100 V. The blotted membranes were blocked with 5% dry milk in 15 TBS buffer and then probed for 2 hr with anti-PARP antibody C2-10 (Trevigen, Gaithersburg, MD). After three 5 min washes with TBS-Tween20 (0.05%), the blots were incubated with secondary antibody, anti-mouse HRPO-anti IgG for 1 hr (Amersham Life Science, Arlington Height IL). Antibody binding was visualized by ECL according to manufacturer's instructions (Amersham Life Science, Arlington Heights, IL).

10
15

Tumors in Nude Mice. Tumor cells (5×10^6) were injected into flanks of female athymic HSD nude mice, at 6-8 weeks of age. The tumors were measured with calipers using the National Cancer Institute formula: $V = L \text{ (mm)} \times I^2 \text{ (mm)} / 2$ where L is the largest diameter and I is the smallest diameter of the tumor. When the volume of the nodules had achieved about 100-150 mm³, tumor-bearing mice were assigned randomly for the control or treatment groups (3-5 mice / group).

20

Antitumor Effects of TMZ and MX plus TMZ. Nude mice carrying tumors were received TMZ, MX or MX combined with TMZ daily for a period of 5 days of treatment. Doses used per group were as follows: a) TMZ alone, 120 mg/kg; b) MX alone, 2 mg/kg; c) BG plus TMZ, 30 mg/kg BG + 120 mg/kg; d) MX plus TMZ, 0.2 mg/kg MX + 120 mg/kg TMZ.

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Experimental End Point. Tumor measurements were taken every 3 days. The relative tumor volume (V/V_0) was calculated by dividing the measured tumor volume (V) by the initial tumor volume (V_0) at day 0. Tumor responses were

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quantified by tumor regrowth delay. Tumor growth delays were calculated according to: tumor growth delay = $T_{2X} - C_{2X}$, where T_{2X} and C_{2X} represent the number of days treated and control tumors take to double in size from the day of treatment, respectively.

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Toxicity Evaluation. Toxicity after treatment was evaluated by body weight measurements and peripheral WBC counting. Body weight was measured three times weekly from the first treatment until 2 weeks after the end of treatment. The weight loss was expressed as a percentage of the initial weight (initial weight - lowest weight / initial weight x 100%). Peripheral WBCs were monitored 5 days after the end of treatment compared with WBCs of control mice.

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EXAMPLE 1

Methoxyamine Potentiates Cytotoxicity of TMZ

We have previously reported the comparative cytotoxicity of TMZ and BG in the SW480 and HCT116 cells lines (Liu *et al.* "Mismatch repair mutations override alkyltransferase in conferring resistance to temozolomide but not to 1,3-bis(2-chloroethyl)nitrosourea" *Cancer Res.* 56: 5375-5379, 1996). In order to test whether MX would alter TMZ cytotoxicity, we treated SW480 and HCT116 with 6 mM MX (itself a nontoxic concentration) plus TMZ (0-1500 μ M) for 2 hr, with or without BG to abolish AGT mediated removal of O^6 mG DNA adducts. SW480 cells were moderately resistant to TMZ, with an IC_{50} of 395 μ M which was reduced 14-fold to 28 μ M by BG pretreatment. Greater resistance to TMZ was observed in MH1 defective cells, even after inhibition of AGT by BG (IC_{50} , 950 μ M). In both cell lines, MX potentiated the cytotoxic effect of TMZ with a DMF of 2.4 in SW480 and 3.1 in HCT116 (Fig. 1). In SW480 cells, additive effects of MX and BG were noted (IC_{50} was reduced from 395 μ M to 6 μ M), whereas, with HCT116 cells no effect of BG was seen in the presence of MX (Fig. 1: A. HTC116 cells; B. SW480 cells. (■), TMZ alone; (◆), MX plus TMZ; (●), BG plus TMZ; (▲), BG and MX plus TMZ).

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To further decipher the role of N^3 mA and N^7 mG DNA adducts in the relative absence of O^6 mG, we evaluated the effect of MX on MMS mediated cytotoxicity. NMS is a methylating agent which produces far fewer O^6 mG adducts (0.3%) and a

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greater proportion of N³mA (10%) and N⁷mG adducts (87%) than TMZ (Chou and Talalay "Quantitative analysis of dose-effect relationship: the combined effects of multiple drugs on enzyme inhibitors" In: G. Weber (ed.) *Advances in Enzyme Regulation*, pp. 27-55, New York: Pergamon Press, 1983). The IC₅₀ of MMS was 0.82 mM in SW480 and 1.4 mM in HCT116 cells. This difference is smaller than the difference in the TMZ IC₅₀ between these cell lines, probably because the low level of O⁶mG adducts formed by MMS increases the impact of other DNA adducts. After cells were treated with MMS (0-3 mM) plus 6 mM MX for 1 hr, the IC₅₀ DMFs, compared to MMS alone, were 2.0 in SW480 and 2.3 in HCT116 (Fig. 2: A. HCT116 cells; B. SW480 cells. (■), MMS alone; (◆), MX plus MMS; (●), BG plus MMS; (▲), BG and MX plus MMS). These DMFs were similar to that observed with TMZ. Compared to treatment of SW480 with BG plus TMZ (DMF was 14), BG plus MMS induced less enhancement of cytotoxicity (DMF was 6). When MMS was combined with BG and MX, >10 fold potentiation of cytotoxicity was observed in SW480 cells, whereas no additive effect was seen in HCT116 cells. In SW480 cells, the increased MMS-toxicity by BG suggests that despite the small number of O⁶mG adducts, they contribute to cytotoxicity. In HCT116 cells, increased cytotoxicity was only observed in the combination of TMZ with MX but not with BG, indicating that MX interfered with processing of N⁷mG and N³mA lesions which killed cells. From these data, we infer that MX had equal ability to interrupt BER in these two cell lines.

EXAMPLE 2

Inhibitors of PARP Modulate the Sensitivity of Cells to TMZ

Since inhibitors of PARP may interrupt BER and increase sensitivity to methylating agents, we examined whether inhibitors of PARP sensitize cells to TMZ. Fig. 3a and Fig. 4a ((■), TMZ alone; (◆), BG plus TMZ; (●), PD128763 plus TMZ; (▲), BG and TMZ plus PD128763) display survival after combined treatment of TMZ with PD128763, 3-AB or 6-AN in both SW480 and HCT116 cells. In the SW480 cell line, PD128763 sensitized cells to TMZ with a DMF of 3.3-fold. The combination of PD128763, BG, and TMZ was even more toxic, with a DMF compared to TMZ alone of 36-fold. In HCT116 cells, the DMF for PD128763

and TMZ compared to TMZ alone was 5.0. However, the combination of PD128763, BG and TMZ had no greater effect than PD128763 and TMZ, indicating that persistent O⁶mG had no effect on cytotoxicity in this MMR defective cell line. Potentiation of TMZ cytotoxicity was also observed in both cell lines treated with two other PARP inhibitors, 3-AB (Fig. 3b, 4b: (■), TMZ alone; (◆), BG plus TMZ; (●), 6-AN plus TMZ; (▲), BG and TMZ plus 6-AN) and 6-AN (Fig. 4c, 4c: (■), TMZ alone; (◆), BG plus TMZ; (●), 3-AB plus TMZ; (▲), BG and TMZ plus 3-AB). Although the specific activity of these agents varied considerably, the DMFs were similar, 3-4-fold, for both 3-AB and 6-AN.

EXAMPLE 3

Synergistic Interaction Between TMZ and MX or PD128763

We investigated the nature of the reaction between TMZ and MX in the TMZ-resistant cell line, HCT116. These cells were incubated in the presence of a range of concentrations of TMZ (75-750 μ M), of MX (1.5-15.0 mM), or constant molar ratio mixture of TMZ and MX (1:20) for 2 hr. HCT116 cells were also exposed to TMZ and PD128763 (62.5-625 μ M) alone and to a combination of (1:0.83) for 2 hr to analyze synergism. As shown in Fig. 5A and 5B, synergistic interaction ($CI \ll 1$, $p < 0.001$) was found in both SW480 cells and HCT116 cells using the combination of TMZ with either MX or PD 128763. These results also showed a marked synergism at high Fa values. Since MDF for these combinations was similar to the observed with SW480 cells, we conclude that BER inhibitor synergizes methylating agent cytotoxicity in both MMR deficient and proficient colon cancer cells.

EXAMPLE 4

Effect of BER Inhibitors on BCNU Cytotoxicity

To test whether MX is also able to sensitize colon cancer cells to chloroethylating agents, these two cell lines were pretreated with 6 mM MX for 2 hr followed by BCNU. No enhancement of BCNU cytotoxicity by MX was observed (Fig. 6); the BCNU IC₅₀ was 45 μ M in HCT116 cells (Fig. 6A: (■), BCNU alone;

(◆) MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU) and 27-29 μM in SW480 cells, respectively, treated with BCNU alone or BCNU plus MX (Fig 6B: (■), BCNU alone; (◆), MX plus BCNU; (●), BG plus BCNU; (▲), BG plus MX plus BCNU). A greater sensitization to BCNU was observed in these two cell lines when cells were treated with MX plus BG and BCNU, in which the BCNU IC_{50} for both cell lines was 5 μM . However, most of the effect was potentiated due to BG, which increased BCNU cytotoxicity by 3-4 fold. As shown in Fig. 7, no sensitization to BCNU cytotoxicity was seen after treatment with PD 128763 or BG and PD 128763 ((■), BCNU alone; (◆), PD 128763 plus BCNU; (●), BG plus BCNU; (▲), BG and PD128763 plus BCNU; (▶), BG, PD128763 and MX plus BCNU).

EXAMPLE 5

Effect of Inhibitors of BER on Cell Cycle Distribution and PARP Cleavage

The cell cycle and apoptosis response of SW480 and HCT116 cells was examined at various times after treatment with TMZ (300 μM) alone or with either MX (6 mM), PD128763 (100 μM) or BG (25 μM). After treatment, cells were divided into two aliquots for analysis of cell cycle / apoptosis on days 1 and 3, and (see below) for detection of PARP cleavage. Cell cycle distribution was measured by flow cytometry according to DNA content and estimation of the duration of G1, S and G2/M was based on untreated, exponentially growing, asynchronous cells. MX and PD128763 alone did not affect the distribution of the cell cycle in these two cell lines (data not shown). At 24 hr, 75 - 90% of SW480 cells accumulated in S and G2 after treatment with TMZ alone and this G2/S phase arrest was more pronounced in cells pretreated with either MX or PD128763 (Fig. 8A). S/G2 arrest was still present at 3 days in cells treated with the combination of MX or PD128763 and TMZ (in both instances, 13 - 20% of the cells were apoptotic). In SW480 cells treated with TMZ alone, the G2/S block was less obvious at day 3 with only 8% of the cells showing evidence of apoptosis. In contrast, HCT116 cells had a normal cell cycle distribution after treatment with TMZ alone and no effect was seen with BG and

TMZ. However, accumulation in S phase was observed (Fig 8B) 24 hr after treatment with PD128763 plus TMZ. At 72 hr, HCT116 cells had moved through S phase and thereafter, a significant portion of cells (90%) remained arrested in G2 with apoptosis present in 1 % of the cells. A similar but less striking result was observed with MX and TMZ in HCT116 cells. By 72 hr, 60% of cells were still arrested in S and G2 and 10% of the cells were apoptotic.

Finally, as a marker of apoptosis induced cell death, we examined PARP cleavage after cells were treated with these drug combinations at 3 days (Fig. 9). PARP cleavage was observed in SW480 cells after exposure to TMZ alone and TMZ plus BG, but was not seen in HCT116 cells with the same treatment, indicating that the apoptotic process is triggered when O⁶mG lesions are repaired by the MMR system. However, PARP cleavage was detected in MMR wild type and deficient cells treated with TMZ plus either MX or PARP inhibitors.

EXAMPLE 6

Acute Toxicity of MX

Three mice per group were injected (i.p.) with MX at doses of 1, 2, 4, 6, 10 and 15 mg/kg for the test of acute toxicity. MX was lethal at 15 mg/kg causing death within 10 mins. At 10 mg/kg, mice appeared to have decreased motility but recovered one hour after treatment. Lower doses did not appear toxic.

EXAMPLE 7

Antitumor Effect of TMZ or TMZ combined with either BG or MX on SW480 Tumor Xenograft

The response of SW480 xenograft to TMZ alone and combined treatments was shown in Fig 10 and Table 1. At the highest tolerable dose of TMZ, 120 mg/kg, a tumor growth delay of approximately 12 days was noted, indicating that SW480 is sensitive to TMZ. When given 30 mg/kg of BG for one hour prior to 40 mg/kg of TMZ, tumor growth delay was enhanced by BG up to 3-fold. Slightly greater efficacy was seen by combining the BG with high doses of TMZ, but significant weight loss (maximum body weight loss from 26 to 20 g, 23%) and very low leukocyte counts (90% decreased) at day 5 after the last treatment were observed in the mice. In contrast, mice treated with 0.2 mg/kg MX plus 120 mg/kg TMZ had an

immediate cessation of tumor growth for 20 days and very slow regrowth of the tumor with tumor growth delays over 40 days ($p < 0.02$). There was no evidence of toxicity to mice with this combined treatment.

5 Table 1. SW480 xenograft toxicity and tumor response

Group	Doses (mg/kg)	Toxicity (max. BW loss)	Tumor growth delays (Days)
TMZ	120	11% (27±1.9 – 24±2.4 g)	9.3 ± 1.2
MX	2	0%	2.0 ± 0.5
BG+TMZ	30/120	23% (26±1.5 – 20±0.5 g)	10.3 ± 2.4
MX+TMZ	2/120	8% (26±1.5 – 24±1.5 g)	27.0 ± 1.2

EXAMPLE 8

Antitumor Effect of TMZ or TMZ Combined with MX on HCT116

Tumor Xenograft

10 Mice bearing HCT116 xenograft appeared to be relatively sensitive to all treatment. Body weight loss was observed in all treated groups (Table 2). BG plus TMZ caused 28% body weight loss and toxic death (2 of 5 mice) at 5 days after treatment. As shown in Fig. 11, HCT116 xenografts treated either with TMZ or MX alone had similar growth rate and growth delays of tumor treated with 0.5 mg/kg
 15 MX plus 120 mg/kg TMZ were significantly prolonged compared with control tumors ($p < 0.05$).

Table 2. HCT116 xenograft toxicity and tumor response

Group	Doses (mg/kg)	Toxicity (max. BW loss)	Tumor growth delays (Days)
TMZ	120	20.8% (24±1.3 – 19±1.7 g)	2.5 ± 0.8

MX	2	8.3% (24±1.6 – 22±1.4 g)	2.2 ± 0.4
BG+TMZ	30/120	28.0% (25±0.8 – 18±0.5 g)*	NA
MX+TMZ	2/120	19.2% (26±1.3 – 21±0.4 g)	17.0 ± 1.2

* toxic death: 2/5 mice

NA, did not analyze

EXAMPLE 9

5 While an understanding of the underlying mechanism is not necessary to the practice of the invention, and with the understanding that the invention is not to be limited to any particular mechanism, MX potentiates the antitumor effect of methylating agents such as temozolomide (TMZ) both *in vitro* and in human tumor xenograft models. A nonlimiting hypothesis for the underlying mechanism is that the potentiation is mediated through the known ability of MX to bind
10 apurinic/pyrimidinic (AP) sites in DNA. Thus, in this model, one test of which is provided in the instant example, MX would be hypothesized to be useful in chemotherapy against cancer, and to enhance the therapeutic efficacy of a broad class of agents that produce damaged DNA to generate AP sites, thereby interrupting
15 AP-endonuclease (APE) in the base excision repair pathway.

This example shows that the combination of methoxyamine (MX) and 1,3-bis (chloroethyl) 2-nitrosourea (BCNU, a crosslinking agent known to produce AP sites) is effective in the treatment of cancer, using human tumor xenografts grown in nude mice.

20 Tumor cells (5×10^6) were injected into both flanks of female athymic nude mice, at 6-8 weeks of age. The tumors were measured with calipers using the National Cancer Institute formula: $V = L \text{ (mm)} \times I^2 \text{ (mm)} / 2$, where L is the largest diameter and I is the smallest diameter of the tumor. When the volume of tumor nodules has achieved about 100-150 mm³, tumor-bearing mice were assigned
25 randomly for the control or the treatment groups (3-5 mice/group, 2 tumors/mouse).

Nude mice carrying tumor received a single injection (i.p.) of BCNU (30 mg/kg) or MX (2 mg/kg) combined with BCNU.

Tumor measurements were taken every 3 days. The relative tumor volume (V/V_0) was calculated by dividing the measured tumor volume (V) by the initial tumor volume (V_0) at day 0. Tumor responses were quantified by tumor regrowth delay. Tumor growth delays were calculated according to the following formula:
5 tumor growth delay = $T_{2x} - C_{2x}$, where T_{2x} , and C_{2x} represent the number of days treated and control tumors take to double in size from the day of treatment, respectively.

The human colon cancer cell lines HCT116 (hMLH mut., p53 wt and AGT expressing) and HCT116-Ch3 (having restored MLH1 activity) were inoculated into
10 nude mice at 6-8 weeks of age. At the tumor sizes indicated above, the mice received a single i.p. injection, also as described above.

The effects on HCT116 tumors are depicted in Figure 12A. The effects on HCT116-Ch3 are depicted in Figure 12B. At a dose of 30 mg/kg, BCNU alone had a very mild effect on tumor growth. Combined administration of MX and BCNU
15 produced significant tumor growth inhibition. Tumor growth delays ($T_{2x}-C_{2x}$) were 14-16 days in HCT116 tumors (Figure 12A) and HCT116-Ch3 (Figure 12B) respectively. A similar result was also observed in SW480 tumor (MMR wt, p53 mutant and AGT expressing). Tumor growth delays were 25 days for the SW480 tumors (Figure 12C).

20 Importantly, no systemic toxicity was noted in this combined treatment. In contrast, BCNU (at a dose of 25 mg/kg) combined with 06-benzylguanine (30 mg/kg), an AGT inhibitor being tested in clinical trials, caused toxic death at a 100% rate.

In *in vitro* studies, the inventors observed that cells exposed to MX
25 combined with BCNU had increased DNA strand breaks (both single strand breaks (SSB) and double strand breaks (DSB) measured by comet assay and pulsed field gel) compared to BCNU alone in these two colon cell lines.

The potentiation of the BCNU-antitumor effect by MX suggests that the impact of the base excision repair pathway is important for BCNU-induced
30 cytotoxicity and that MX-adducted AP sites efficiently interrupt BER leading to cell death. Again, this is only a hypothesis, and the invention is not to be limited to any

particular mechanism, and the practice of the invention does not require any knowledge of the underlying mechanism.

EXAMPLE 10

This example shows the combination of MX and an oxidizing agent for the treatment of cancer.

Again, while the present invention is not limited to any mechanism, BER comprises a ubiquitous series of biochemical pathways for the removal of oxidative damage to the nitrogenous bases in DNA. Initiating proteins of the BER are DNA glycosylases, which hydrolyze the N-glycosylic bond between the deoxyribose sugar moiety and the DNA base, generating AP sites. AP sites can also be generated by spontaneous destabilization of N-glycosyl bonds, particularly after oxidative damage to the bases. The AP sites are the target for MX, resulting in the disruption of BER.

In this experiment, a growth inhibition assay was carried out with H₂O₂, an oxidizing agent that mimics the effect of chemotherapeutic agents and radiation. Cells in the exponential phase were plated (1 x 10⁴) in 24-well tissue culture plates. After attachment overnight, the cells were incubated with hydrogen peroxide (H₂O₂) (0-150 μM) or MX (6 mM) plus H₂O₂ for 1 hr. Five days after treatment, the media was discarded, and 100 μl of MTT (5 mg/ml in PBS) was added in 1 ml fresh media to each well for 4 hrs at 37 °C. The MTT solution was then removed, and the formazan crystals were dissolved in 200 μl/well of DMSO for 20 min and changes in absorbance were determined at A_{540nm}. Percentage growth inhibition was calculated by comparison of the A_{540nm} reading from treated versus control cells. Drug concentrations that resulted in an IC₅₀ were determined from the plots of percentage growth inhibition versus the logarithm of the drug concentration.

As shown in Figure 13, MX results in significantly increased sensitivity of colon cancer cells, HCT116, to killing induced by H₂O₂.

This result indicates that MX potentially increases the therapeutic efficacy of a class of anticancer drugs that exert cytotoxicity mediated by oxidative damage in DNA, including but not limited to bleomycin, adriamycin and gamma radiation.

EXAMPLE 11

This example is directed to combinations of MX with members of a group of agents including, but not limited to hypoxanthine, 5-FU, uracil, IUdR, bleomycin, adriamycin and gamma radiation.

The agents contemplated for combinations with MX in this example are agents which incorporate into DNA and are repaired by BER to form AP sites. Cell-growth inhibition was carried out with HCT116 cells to test MX-potentiated cytotoxicity of iododeoxyuridine (IUdR), which has been recognized as a radiosensitizing agent since the early 1960's. As shown in Table 3, MX increases the killing effect of IUdR (or MX decreases the IC50 of >>60 μ M for IUdR alone to 35 μ M) in MMR deficient HCT116 cells. This suggests that MX and IUdR combined with radiation therapy will further increase the radiosensitivity of tumor cells.

These experiments demonstrate that a significant enhancement of antitumor effect of TMZ by MX or PARP inhibitors in human colon cancer xenographs with mismatch repair proficiency and deficiency. Therefore, from the above it should be clear that the present invention provides a wide variety of ways to 1) screen for compounds that can potentiate agents useful in the treatment of cancer, 2) provide model systems for the study of cancer treatment by agents that modulate DNA repair mechanisms and, 3) provide treatments for various cancers.

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Table 3

	IUdR	IUdR + MX (6 mM)
IC 50 (μ M)	>>60	35

EXAMPLE 12

We observed that MX synergistically increased TMZ-induced cytotoxicity in colon cancer cell lines in both MMR proficient and deficient cells. We hypothesized that TMZ-induced N3mA and N7mG DNA adducts generate AP sites as the major targets for MX-potential, and thus that the formation and persistence of these DNA adducts (previous study showed that N7mG half life was 24-48 hrs) become

the potential lesions causing cell death in the presence of MX. To test this hypothesis, we treated cells with TMZ+MX (25 mM) for 2 hrs and after washing out drugs, we added low concentrations of MX (2.5 mM) over a 72-hr period in colon cancer cell lines. Cell lines used in this study include SW480 (MMR wt, p53 mut), HCT116 (hMLH1 mut, p53 wt) and HCT15 (hMSH6 mut, p53 mut). We measured AP sites using ARP reagent, noting that ARP and MX compete for binding to AP sites. In SW480 cells, AP sites peaked at 2 h after exposure to TMZ (375 μ M, IC₅₀ for this cell line) alone and half maximal amounts of AP sites remained at 72 hr. In contrast, lower levels of AP sites were detected (20% of AP sites generated by TMZ alone) over a period of 72 h when cells were treated with MX and TMZ, because MX-adducted AP sites were not recognized by ARP. Thus 80% of AP sites were MX bound. Prolonged exposure to MX apparently results in more MX-AP sites during repair of N7mG. Moreover, after treatment, 50% of cells accumulate in S phase resulting in increased DNA double strand breaks (DSB) detected by pulsed field gel assay. S phase arrest indicates that DNA replication was blocked by DNA damage, as a consequence of MX-adducted AP sites. The combination of MX and TMZ also induced topoisomerase IIa, whereas topo I levels remained consistent, similar to the effect of the known topo II inhibitor, etoposide. This suggests that MX-AP sites form complexes with topo I and II. It is likely that cytotoxicity elicited by both topo I/II inhibitors and MX-AP sites is due to DNA cleavable complexes, and is thus dependent on the level of topo I/II expression. To exploit this pathway, we added a topo II inhibitor, etoposide (5 μ M) to the combination of TMZ and MX and found a remarkable increase in DSB. At 72 hr after treatment, this profound degree of DSB was not noted in cells treated with etoposide and TMZ. MX-AP sites (more than AP sites) thus act as a topo II poison and synergistically enhance the effect of a topo II inhibitor to stabilize cleavable complexes with DNA.

EXAMPLE 13

As demonstrated above, both O⁶benzylguanine (BG) and methoxyamine (MX) potentiate the antitumor effect of methylating agents such as temozolomide (TMZ) in human tumor xenografts in nude mice. We observed less myelosuppression and less weight loss using TMZ and MX than using TMZ and BG. These results suggested that MX may not potentiate TMZ cytotoxicity in

hematopoietic cells. To test this hypothesis, we conducted *in vitro* experiments comparing the sensitivity to BG+TMZ and MX+TMZ in colon cancer cell lines with that of mouse bone marrow and human bone marrow cells. We found that BG potentiated TMZ-cytotoxicity up to 14 fold in SW480 tumor cells and 10 fold in human and mouse bone marrow cells respectively. These results indicate the sensitization of BG to TMZ is similar in the two cell types and results from the inactivation of AGT. MX increased TMZ killing by 4 fold in SW480 cells. However, no potentiation of TMZ cytotoxicity was observed with addition of MX in either human CD34+ cells or murine bone marrow progenitor cells when analyzed by colony forming assay. Topoisomerase (Topo) I and Topo II recognize spontaneous AP sites and form stable cleavable complexes. We hypothesized that AP sites formed during repair of TMZ-induced N7mG and N3mA adducts would also be recognized by Topo I and Topo II. However since Base Excision Repair [BER] is normally quite efficient, we further hypothesized that the impact of Topo I and Topo II would be most important after MX inhibition of BER; increased unrepaired MX adducted AP sites should increase Topo mediated strand breaks and apoptosis. Therefore, potentiation of TMZ cytotoxicity by MX would be dependent on Topo I and II. We found that levels of topoisomerases I and II were much higher (> 20-fold) in human tumor cell lines than in either human CD34+ or murine bone marrow progenitor cells. Thus, we propose that MX does not potentiate TMZ toxicity in hematopoietic progenitors because these cells express low levels of Topo I and II. Lack of Topo I and II results in less cleavable complex formation, fewer double strand DNA breaks, and less apoptosis. We predict a better therapeutic index for the combination of MX/TMZ than observed with BG/TMZ because of the relative overexpression of topoisomerases I and II in malignant cells compared to marrow progenitors.

EXAMPLE 14

The results that methoxyamine (MX) potentiated the cytotoxic effect of temozolomide (TMZ) and BCNU indicated that molecules targeting abasic lesions in DNA are important approach to improve chemotherapeutic efficacy of alkylating agents. On the basis of a similar molecular reaction to MX, (e.g., a primary amine with the carbonyl group of the abasic site), we tested whether a compound

(Compound A) would have the ability to bind to AP site generated by TMZ and enhance the killing effect of TMZ. A has a MX-like structure to bind to AP site and also has fluorescence molecules to provide a signal for direct detection of modified AP sites using cellular image. We found that this compound was able to bind to AP sites assayed with aldehyde-reactive probe (ARP) that competitively binds to the aldehyde group of AP site. A-modified AP sites were refractory to cleavage of APE, suggesting that it can block base excision repair (BER) pathway by inhibiting repair of AP sites. Cellular image showed 30% of cells with fluorescent signal that was located in nuclear at 2 hr after treatments with TMZ plus A. In contrast, cells treated with A alone had no visualized fluorescence signal. This data suggest that A specifically binds to AP sites in DNA. In *in vitro* study, we observed A at non-toxic concentrations (1 mM) sensitized cells to TMZ-cytotoxicity 2-3-fold. In a xenograft model, A (2 mg/kg) combined with TMZ (80 mg/kg) enhanced anti-tumor effect in HCT 116 and DLD1, two xenograft tumors with very high resistance to TMZ. Tumor growth delays were 13 days in HCT 116 and 12 days in DLD1 treated with the combination of A and TMZ over TMZ alone. Thus, A has a similar effect to MX on interruption of BER and can potentiate chemotherapeutic agents such as alkylating agents. Results are depicted in Figures 14-16.

EXAMPLE 15

The experiments above suggest that MX synergistically increased TMZ-induced cytotoxicity in colon cancer with either MMR proficiency or deficiency as a result of TMZ- AP sites generated from induced N3mA and N7mG DNA adducts. AP sites are the major targets being responsible for MX-induced potentiation of alkylating therapeutic agents.

In vivo studies showed that the treatment with MX and TMZ induced extremely high frequencies of chromosomal aberration (present in over 90% of chromosomes) in tumor xenografts after three daily treatments, including aneuploidy, sister chromosome exchange, and severe DNA breakage. The single strand and double strand DNA breaks (Fig. 4) are visible at metaphase. It is well known that topoisomerases are essential enzymes that interconvert topological forms of DNA for transcription, DNA replication, and chromosome segregation,

and it has been demonstrated that AP sites, when located within a topoisomerase I/II DNA cleavage site, act as topoisomerase poisons and stimulate enzyme-mediated DNA scission. Therefore, these results suggest that topoisomerases may be involved in the induction of DNA aberration induced by TMZ and MX. We
5 hypothesize that MX-AP sites act as topo II poisons to abort cleavage and ligation reaction of topo II (higher levels of endogenous topoisomerase II or) and induced levels of topoisomerase II will promote topo II mediated-DNA cleavage resulting in accumulation of DNA double-strand breaks. To test this hypothesis, we developed a combined treatment with TMZ, MX, and a topoII inhibitor, etoposide,
10 in colon cancer cell lines.

Methods:

Analysis of the effect of AP site or MX-AP site on topoisomerase II-mediated DNA cleavage. A position-specific apurinic site was incorporated by replacing a single nucleoside with deoxyuridine at a topoisomerase II cleavage site
15 and then removing the uracil base with uracil-DNA glycosylase, generating AP sites that further are incubated with MX to produce MX-AP sites (Fig 17).

Measurement of DNA double-strand breaks by pulsed-field gel electrophoresis (PFGE). A PFGE procedure was used to visualize induction of
20 DSBs. Log-phase cultures were exposed to the DNA-damaging agents for 15 min at 37 °C, then cells were harvested by scraping into ice-cold PBS. Cells were centrifuged and washed twice with ice-cold PBS. The cells were resuspended in an appropriate volume of PBS, and an equal volume of 1.5% low melting point agarose in TAE buffer to yield 5×10^6 cells per 75 μ l plug. The plugs were allowed
25 to solidify on ice, then were incubated at 50 °C overnight in approximately 5 ml of a lysis solution of 1% sarcosyl, 100 mM EDTA, 10 mM Tris-HCl, pH 7.5, and 100 μ g/ml of proteinase K. The following day, the lysis solution was replaced with TE buffer (10 mM Tris-HCl, pH 7.5 with 50 mM EDTA). Plugs were inserted into the wells of a 0.7% agarose gel. The gel was run using a clamped

homogeneous electric field (CHEF) Mapper XA system (BioRad) using the 180° field inversion gel electrophoresis (FIGE) program; at electrophoresis conditions that can achieve the best sensitivity in detection of DNA DSB and good resolution in the megabase size region. After electrophoresis, the gel was stained with
5 ethidium bromide to visualize DNA.

Median Effect Analysis (MEA). MEA was used to determine the dose-response interactions between Docetaxel and Rebecomycine. Drugs were combined at the ratio of the IC₅₀ values for either Docetaxel alone or Rebecomycine alone as
10 determined by survival/concentration curves. The combination was compared to the cytotoxicity of each drug alone in every experiment. The combination index (CI) was determined from survival fractions at increasing levels of cell killing, using analysis of multiple drug interaction program (Biosoft, Cambridge, United
15 Kingdom) developed based on a method of Chou and Talalay. CI values of less than or greater than 1 indicate synergy and antagonism, respectively, whereas a CI value of 1 indicates additivity of the drugs.

Results:

First, we examined whether APE has a differential effect between regular AP sites and MX-AP sites located in a position specific site for topo II cleavage.
20 Results show that APE is able to cleave regular AP sites rather than MX-bound AP sites (Fig. 18), however, both AP and MX-AP sites are cleaved by topoisomerase II, indicating MX-AP sites are able to stimulate topoisomerase II-mediated DNA cleavage. The cleavage is also observed after incubation of DNA substrates with nuclear extract of colon cancer cells, even having pretreatment with etoposide,
25 indicating that MX-AP site-stimulated Topo II cleavage is independent of the presence of etoposide (Fig. 19).

We predicted that after treatment with MX and TMZ to generate MX-AP sites, the DNA lesion is able to trap topo II in cleavage complex, whereas etoposide interacts with topo II to form Topo II-DNA cleavage complexes. We

thus suspected that they all convert topo II into a physiological toxin that creates DNA double strand breaks, thus, synergistically inducing cell death.

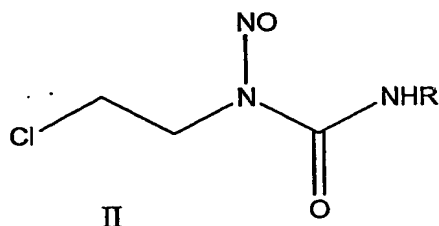
To exploit this pathway, we added a topo II inhibitor, etoposide (5 μ M), to the combination of TMZ and MX and found a remarkable increase in DSB. Even at 72 hr after treatment, this profound degree of DSB was noted in cells treated with etoposide, TMZ, and MX (Fig. 20).

Colony survival assay shows that the combination treatment increased the cell-killing effect (Fig. 21A), and evidenced a synergistic interaction between TMZ, MX, and etoposide (Fig. 21B) when analyzed with multiple drug interaction program (based on the method of Chou and Talalay: combination index values of less than or greater than 1 indicate synergy and antagonism, whereas a combination index value of 1 indicates additivity of the drugs).

In summary, MX-AP sites (to a greater degree than AP sites) act as topo II poisons and synergistically enhance the effect of a topo II inhibitor, etoposide, by stabilizing cleavable complexes with DNA and increasing DNA double-strand breaks, leading to increased cell death.

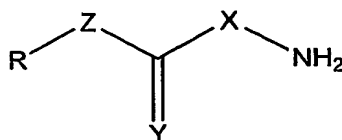
We claim:

1. A formulation comprising an anticancer agent and a base excision repair (BER) inhibitor admixed with pharmaceutically acceptable excipient, wherein the anticancer agent induces formation of AP sites.
- 5 2. The formulation of claim 1, wherein said anticancer agent is selected from a DNA oxidizing agent, ultraviolet radiation, a DNA intercalating agent, a radiosensitizing agent, a cross-linking agent, and an alkylating agent.
3. The formulation of claim 2, wherein said anticancer agent is a cross-linking agent.
- 10 4. The formulation of claim 3, wherein said a cross-linking agent is a mustine having the structure of formula II:



wherein R is an optionally substituted hydrocarbon substituent.

- 15 5. The formulation of claim 4, wherein said mustine is BCNU.
6. The formulation of any one of claims 2, 4, or 5, wherein said BER inhibitor is an AP endonuclease inhibitor.
7. The formulation of claim 6, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

wherein X is O or NH,

Y is O, S, or NH,
Z is absent or represents O, S, or NH, and
R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

5

8. The formulation of claim 7, wherein said AP endonuclease inhibitor is compound A.

9. The formulation of claim 6, wherein said formulation further comprises a topoisomerase inhibitor.

10 10. The formulation of claim 9, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

11. The formulation of claim 10, wherein the topoisomerase II inhibitor is etoposide.

12. The formulation of any one of claims 2, 4, or 5, wherein said BER inhibitor
15 is a PARP inhibitor.

13. The formulation of claim 12, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

14. The formulation of claim 13, wherein said formulation further comprises a topoisomerase inhibitor.

20 15. The formulation of claim 14, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

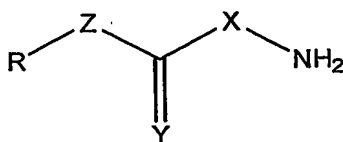
16. The formulation of claim 15, wherein the topoisomerase II inhibitor is etoposide.

25 17. The formulation of claim 2, wherein said anticancer agent is an alkylating agent.

18. The formulation of claim 17, wherein said alkylating agent is temozolomide.

19. The formulation of any one of claims 17 or 18, wherein said BER inhibitor is an AP endonuclease inhibitor.

20. The formulation of claim 19, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

10 R represents a hydrogen or a hydrocarbon moiety, and pharmaceutically acceptable salts thereof.

21. The formulation of claim 20, wherein said AP endonuclease inhibitor is compound A.

22. The formulation of claim 20, wherein said formulation further comprises a
15 topoisomerase inhibitor.

23. The formulation of claim 22, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

24. The formulation of claim 23, wherein the topoisomerase II inhibitor is etoposide.

20 25. The formulation of claim 2, wherein said anticancer agent is a DNA oxidizing agent.

26. The formulation of claim 25, wherein the DNA oxidizing agent is selected from hydrogen peroxide, bleomycin, and adriamycin.

27. The formulation of claim 2, wherein said anticancer agent is a
25 radiosensitizing agent.

28. The formulation of claim 27, wherein said radiosensitizing agent is IUdR.

29. The formulation of claim 2, wherein said anticancer agent is selected from uracil, hypoxanthine, and 5-FU.

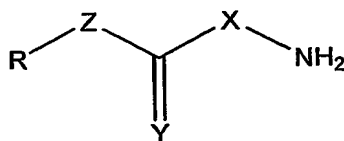
30. The formulation of claim 2, wherein said anticancer agent is ultraviolet
5 radiation.

31. The formulation of claim 30, wherein said ultraviolet radiation is gamma irradiation.

32. The formulation of claim 2, wherein said anticancer agent is selected from temozolomide, IUdR, hydrogen peroxide, bleomycin, adriamycin, uracil,
10 hypoxanthine, and 5-FU.

33. The formulation of claim 32, wherein said BER inhibitor is an AP endonuclease inhibitor.

34. The formulation of claim 33, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



15

Formula I

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

20 R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

35. The formulation of any one of claims 17 or 18, wherein said BER inhibitor is a PARP inhibitor.

36. The formulation of claim 35, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

37. The formulation of claim 35, wherein said formulation further comprises a topoisomerase inhibitor.

5 38. The formulation of claim 37, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

39. The formulation of claim 38, wherein the topoisomerase II inhibitor is etoposide.

10 40. The formulation of claim 1, wherein a dose of the formulation comprises an amount of the anticancer agent that is subtherapeutic when administered in the absence of the base excision repair inhibitor.

41. The formulation of claim 1, wherein said base excision repair inhibitor is selected from an AP endonuclease inhibitor, a DNA glycosylase inhibitor, a DNA polymerase inhibitor, a PARP inhibitor, and a DNA ligase inhibitor.

15 42. The formulation of claim 41, wherein said base excision repair inhibitor is an AP endonuclease inhibitor.

43. The formulation of claim 41, wherein said base excision repair inhibitor is a PARP inhibitor.

20 44. The formulation of claim 43, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

45. The formulation of claim 41, wherein said BER inhibitor is an inhibitor of DNA polymerase.

46. The formulation of claim 45, wherein said inhibitor of DNA polymerase inhibits DNA polymerase β , γ , or ϵ .

47. The formulation of claim 41, wherein said base excision repair inhibitor is a DNA ligase inhibitor.

48. The formulation of claim 47, wherein said DNA ligase inhibitor inhibits the action of DNA ligase I or DNA ligase II.

5 49. The formulation of any one of claims 7, 10, 13, 16, 20, 24 or 32 which is administered to a patient diagnosed with cancer.

50. The formulation of claim 49, wherein said formulation is administered orally.

10 51. The formulation of claim 49, wherein said formulation is administered intravenously.

52. The formulation of claim 49, wherein the cancer is selected from carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.

15 53. The formulation of claim 1, wherein said formulation further comprises a DNA alkyltransferase inhibitor.

54. The formulation of claim 53, wherein said DNA alkyltransferase inhibitor is BG.

55. The formulation of claim 1, wherein said formulation further comprises a topoisomerase inhibitor.

20 56. The formulation of claim 55, wherein said topoisomerase inhibitor is a topoisomerase I inhibitor.

57. The formulation of claim 55, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

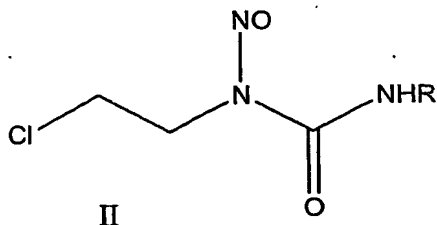
25 58. The formulation of claim 57, wherein the topoisomerase II inhibitor is etoposide.

59. A method for potentiating a therapeutic effect of an anticancer agent that induces formation of AP sites, comprising administering a base excision repair inhibitor, whereby the base excision repair inhibitor potentiates the effect of the anticancer agent.

5 60. The method of claim 59, wherein said anticancer agent is selected from a DNA oxidizing agent, ultraviolet radiation, a DNA intercalating agent, a radiosensitizing agent, a cross-linking agent, and an alkylating agent.

61. The method of claim 60, wherein said anticancer agent is a cross-linking agent.

10 62. The method of claim 61, wherein said a cross-linking agent is a mustine having the structure of formula II:

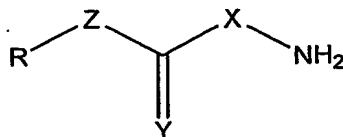


wherein R is an optionally substituted hydrocarbon substituent.

15 63. The method of claim 62, wherein said mustine is BCNU.

64. The method of any one of claims 60, 62, or 63, wherein said BER inhibitor is an AP endonuclease inhibitor.

65. The method of claim 64, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



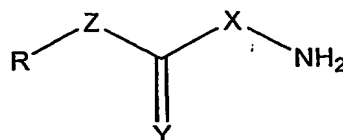
Formula I

wherein X is O or NH,

Y is O, S, or NH,
Z is absent or represents O, S, or NH, and
R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

- 5 66. The method of claim 65, wherein said AP endonuclease inhibitor is compound A.
67. The method of claim 64, wherein said method further comprises administering a topoisomerase inhibitor.
68. The method of claim 67, wherein said topoisomerase inhibitor is a
10 topoisomerase II inhibitor.
69. The method of claim 68, wherein the topoisomerase II inhibitor is etoposide.
70. The method of any one of claims 60, 62, or 63, wherein said BER inhibitor is a PARP inhibitor.
71. The method of claim 70, wherein said PARP inhibitor is selected from
15 PD128763, 3-AB, and 6-AN.
72. The method of claim 71, wherein said method further comprises administering a topoisomerase inhibitor:
73. The method of claim 72, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
- 20 74. The method of claim 73, wherein the topoisomerase II inhibitor is etoposide.
75. The method of claim 60, wherein said anticancer agent is an alkylating agent.
76. The method of claim 75, wherein said alkylating agent is temozolomide.
77. The method of any one of claims 75 or 76, wherein said BER inhibitor is an AP endonuclease inhibitor.

78. The method of claim 77, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

- 5 wherein X is O or NH,
Y is O, S, or NH,
Z is absent or represents O, S, or NH, and
R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.
- 10 79. The method of claim 78, wherein said AP endonuclease inhibitor is compound A.
80. The method of claim 78, wherein said method further comprises administering a topoisomerase inhibitor.
81. The method of claim 80, wherein said topoisomerase inhibitor is a
15 topoisomerase II inhibitor.
82. The method of claim 81, wherein the topoisomerase II inhibitor is etoposide.
83. The method of claim 60, wherein said anticancer agent is a DNA oxidizing agent.
84. The method of claim 83, wherein the DNA oxidizing agent is selected from
20 hydrogen peroxide, bleomycin, and adriamycin.
85. The method of claim 60, wherein said anticancer agent is a radiosensitizing agent.
86. The method of claim 85, wherein said radiosensitizing agent is IUdR.

87. The method of claim 60, wherein said anticancer agent is selected from uracil, hypoxanthine, and 5-FU.

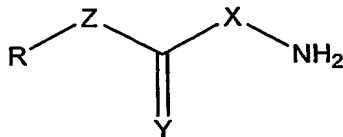
88. The method of claim 60, wherein said anticancer agent is ultraviolet radiation.

5 89. The method of claim 88, wherein said ultraviolet radiation is gamma irradiation.

90. The method of claim 60, wherein said anticancer agent is selected from temozolomide, IUdR, hydrogen peroxide, bleomycin, adriamycin, uracil, hypoxanthine, and 5-FU.

10 91. The method of claim 90, wherein said BER inhibitor is an AP endonuclease inhibitor.

92. The method of claim 91, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

15

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

R represents a hydrogen or a hydrocarbon moiety,

20 and pharmaceutically acceptable salts thereof.

93. The method of any one of claims 75 or 76, wherein said BER inhibitor is a PARP inhibitor.

94. The method of claim 93, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

95. The method of claim 93, wherein said method further comprises administering a topoisomerase inhibitor.

96. The method of claim 95, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

5 97. The method of claim 96, wherein the topoisomerase II inhibitor is etoposide.

98. The method of claim 59, wherein the amount of anticancer agent is subtherapeutic when administered in the absence of the base excision repair inhibitor.

99. The method of claim 59, wherein said base excision repair inhibitor is selected from an AP endonuclease inhibitor, a DNA glycosylase inhibitor, a DNA polymerase inhibitor, a PARP inhibitor, and a DNA ligase inhibitor.

100. The method of claim 99, wherein said base excision repair inhibitor is an AP endonuclease inhibitor.

101. The method of claim 99, wherein said base excision repair inhibitor is a PARP inhibitor.

102. The method of claim 101, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

103. The method of claim 99, wherein said BER inhibitor is an inhibitor of DNA polymerase.

20 104. The method of claim 103, wherein said inhibitor of DNA polymerase inhibits DNA polymerase β , γ , or ϵ .

105. The method of claim 99, wherein said base excision repair inhibitor is a DNA ligase inhibitor.

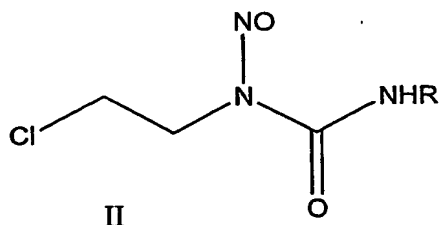
25 106. The method of claim 105, wherein said DNA ligase inhibitor inhibits the action of DNA ligase I or DNA ligase II.

107. The method of any one of claims 65, 68, 71, 74, 78, 82 or 90, wherein the patient is diagnosed with cancer.
108. The method of claim 107, wherein said base excision repair inhibitor is administered orally.
- 5 109. The method of claim 107, wherein said base excision repair inhibitor is administered intravenously.
110. The method of claim 107, wherein the cancer is selected from carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.
- 10 111. The method of claim 59, wherein said method further comprises administering a DNA alkyltransferase inhibitor.
112. The method of claim 111, wherein said DNA alkyltransferase inhibitor is BG.
113. The method of claim 59, wherein said method further comprises
15 administering a topoisomerase inhibitor.
114. The method of claim 113, wherein said topoisomerase inhibitor is a topoisomerase I inhibitor.
115. The method of claim 113, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
- 20 116. The method of claim 115, wherein the topoisomerase II inhibitor is etoposide.
117. A kit comprising a first pharmaceutical preparation comprising an anticancer agent that induces formation of AP sites, a second pharmaceutical preparation comprising a base excision repair inhibitor, and instructions for administering the
25 first and second pharmaceutical preparations to a patient for the treatment of cancer.

118. The kit of claim 117, wherein said anticancer agent is selected from a DNA oxidizing agent, ultraviolet radiation, a DNA intercalating agent, a radiosensitizing agent, a cross-linking agent, and an alkylating agent.

119. The kit of claim 118, wherein said anticancer agent is a cross-linking agent.

5 120. The kit of claim 119, wherein said a cross-linking agent is a mustine having the structure of formula II:

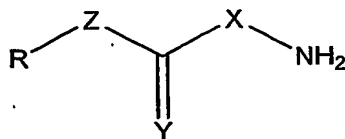


wherein R is an optionally substituted hydrocarbon substituent.

10 121. The kit of claim 120, wherein said mustine is BCNU.

122. The kit of any one of claims 118, 120, or 121, wherein said BER inhibitor is an AP endonuclease inhibitor.

123. The kit of claim 122, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

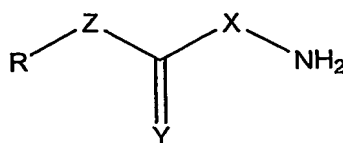
wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

15 20 R represents a hydrogen or a hydrocarbon moiety, and pharmaceutically acceptable salts thereof.

124. The kit of claim 123, wherein said AP endonuclease inhibitor is compound A.
125. The kit of claim 122, wherein said kit further comprises a topoisomerase inhibitor.
- 5 126. The kit of claim 125, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
127. The kit of claim 126, wherein the topoisomerase II inhibitor is etoposide.
128. The kit of any one of claims 118, 120, or 121, wherein said BER inhibitor is a PARP inhibitor.
- 10 129. The kit of claim 128, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.
130. The kit of claim 129, wherein said kit further comprises a topoisomerase inhibitor.
131. The kit of claim 130, wherein said topoisomerase inhibitor is a
15 topoisomerase II inhibitor.
132. The kit of claim 131, wherein the topoisomerase II inhibitor is etoposide.
133. The kit of claim 118, wherein said anticancer agent is an alkylating agent.
134. The kit of claim 133, wherein said alkylating agent is temozolomide.
135. The kit of any one of claims 133 or 134, wherein said BER inhibitor is an AP
20 endonuclease inhibitor.
136. The kit of claim 135, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

wherein X is O or NH,

Y is O, S, or NH,

- 5 Z is absent or represents O, S, or NH, and
R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

137. The kit of claim 136, wherein said AP endonuclease inhibitor is compound A.

- 10 138. The kit of claim 136, wherein said kit further comprises a topoisomerase inhibitor.

139. The kit of claim 138, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

140. The kit of claim 139, wherein the topoisomerase II inhibitor is etoposide.

- 15 141. The kit of claim 118, wherein said anticancer agent is a DNA oxidizing agent.

142. The kit of claim 141, wherein the DNA oxidizing agent is selected from hydrogen peroxide, bleomycin, and adriamycin.

- 20 143. The kit of claim 118, wherein said anticancer agent is a radiosensitizing agent.

144. The kit of claim 143, wherein said radiosensitizing agent is IUdR.

145. The kit of claim 118, wherein said anticancer agent is selected from uracil, hypoxanthine, and 5-FU.

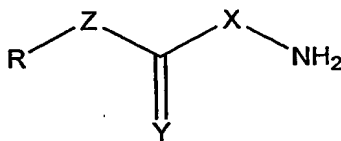
146. The kit of claim 118, wherein said anticancer agent is ultraviolet radiation.

147. The kit of claim 146, wherein said ultraviolet radiation is gamma irradiation.

148. The kit of claim 118, wherein said anticancer agent is selected from temozolomide, IUdR, hydrogen peroxide, bleomycin, adriamycin, uracil, hypoxanthine, and 5-FU.

5 149. The kit of claim 148, wherein said BER inhibitor is an AP endonuclease inhibitor.

150. The kit of claim 149, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



10 Formula I

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

R represents a hydrogen or a hydrocarbon moiety.

15 and pharmaceutically acceptable salts thereof.

151. The kit of any one of claims 133 or 134, wherein said BER inhibitor is a PARP inhibitor.

152. The kit of claim 151, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

20 153. The kit of claim 151, wherein said kit further comprises a topoisomerase inhibitor.

154. The kit of claim 153, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

155. The kit of claim 154, wherein the topoisomerase II inhibitor is etoposide.

156. The kit of claim 117, wherein the amount of the anticancer agent is subtherapeutic when administered in the absence of the base excision repair inhibitor.
157. The kit of claim 117, wherein said base excision repair inhibitor is selected
5 from an AP endonuclease inhibitor, a DNA glycosylase inhibitor, a DNA polymerase inhibitor, a PARP inhibitor, and a DNA ligase inhibitor.
158. The kit of claim 157, wherein said base excision repair inhibitor is an AP endonuclease inhibitor.
159. The kit of claim 157, wherein said base excision repair inhibitor is a PARP
10 inhibitor.
160. The kit of claim 159, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.
161. The kit of claim 157, wherein said BER inhibitor is an inhibitor of DNA polymerase.
- 15 162. The kit of claim 161, wherein said inhibitor of DNA polymerase inhibits DNA polymerase β , γ , or ϵ .
163. The kit of claim 157, wherein said base excision repair inhibitor is a DNA ligase inhibitor.
164. The kit of claim 163, wherein said DNA ligase inhibitor inhibits the action of
20 DNA ligase I or DNA ligase II.
165. The kit of any one of claims 117, 123, 126, 129, 132, 136, 140 or 148, wherein the cancer is selected from carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.
- 25 166. The kit of claim 117, wherein said kit further comprises a DNA alkyltransferase inhibitor.

167. The kit of claim 166, wherein said DNA alkyltransferase inhibitor is BG.

168. The kit of claim 117, wherein said kit further comprises a topoisomerase inhibitor.

169. The kit of claim 168, wherein said topoisomerase inhibitor is a
5 topoisomerase I inhibitor.

170. The kit of claim 168, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

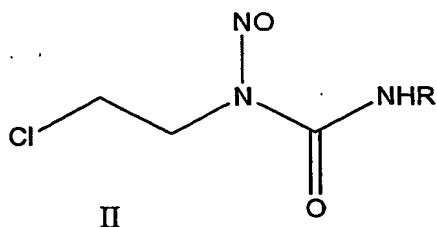
171. The kit of claim 170, wherein the topoisomerase II inhibitor is etoposide.

172. A kit comprising a pharmaceutical preparation comprising a base excision
10 repair inhibitor and instructions for coadministration of the pharmaceutical preparation with an anticancer agent that induces formation of AP sites.

173. The kit of claim 172, wherein said anticancer agent is selected from a DNA oxidizing agent, ultraviolet radiation, a DNA intercalating agent, a radiosensitizing agent, a cross-linking agent, and an alkylating agent.

15 174. The kit of claim 173, wherein said anticancer agent is a cross-linking agent.

175. The kit of claim 174, wherein said a cross-linking agent is a mustine having the structure of formula II:



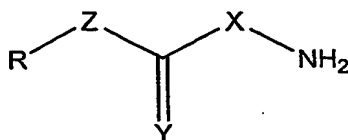
wherein R is an optionally substituted hydrocarbon substituent.

20

176. The kit of claim 175, wherein said mustine is BCNU.

177. The kit of any one of claims 173, 175, or 176, wherein said BER inhibitor is an AP endonuclease inhibitor.

178. The kit of claim 177, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

10 R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.

179. The kit of claim 178, wherein said AP endonuclease inhibitor is compound A.

15 180. The kit of claim 179, wherein said kit further comprises a topoisomerase
inhibitor.

181. The kit of claim 180, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

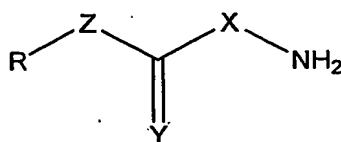
182. The kit of claim 181, wherein the topoisomerase II inhibitor is etoposide.

20 183. The kit of any one of claims 173, 175, or 176, wherein said BER inhibitor is
a PARP inhibitor.

184. The kit of claim 183, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

185. The kit of claim 184, wherein said kit further comprises a topoisomerase inhibitor.

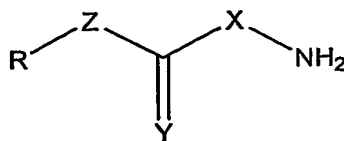
186. The kit of claim 185, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
187. The kit of claim 186, wherein the topoisomerase II inhibitor is etoposide.
188. The kit of claim 173, wherein said anticancer agent is an alkylating agent.
- 5 189. The kit of claim 188, wherein said alkylating agent is temozolomide.
190. The kit of any one of claims 188 or 189, wherein said BER inhibitor is an AP endonuclease inhibitor.
191. The kit of claim 190, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

- wherein X is O or NH,
Y is O, S, or NH,
Z is absent or represents O, S, or NH, and
15 R represents a hydrogen or a hydrocarbon moiety,
and pharmaceutically acceptable salts thereof.
192. The kit of claim 191, wherein said AP endonuclease inhibitor is compound A.
193. The kit of claim 191, wherein said kit further comprises a topoisomerase
20 inhibitor.
194. The kit of claim 193, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
195. The kit of claim 194, wherein the topoisomerase II inhibitor is etoposide.

196. The kit of claim 173, wherein said anticancer agent is a DNA oxidizing agent.
197. The kit of claim 196, wherein the DNA oxidizing agent is selected from hydrogen peroxide, bleomycin, and adriamycin.
- 5 198. The kit of claim 173, wherein said anticancer agent is a radiosensitizing agent.
199. The kit of claim 198, wherein said radiosensitizing agent is IUdR.
200. The kit of claim 173, wherein said anticancer agent is selected from uracil, hypoxanthine, and 5-FU.
- 10 201. The kit of claim 173, wherein said anticancer agent is ultraviolet radiation.
202. The kit of claim 201, wherein said ultraviolet radiation is gamma irradiation.
203. The kit of claim 173, wherein said anticancer agent is selected from temozolomide, IUdR, hydrogen peroxide, bleomycin, adriamycin, uracil, hypoxanthine, and 5-FU.
- 15 204. The kit of claim 203, wherein said BER inhibitor is an AP endonuclease inhibitor.
205. The kit of claim 204, wherein said AP endonuclease inhibitor is selected from methoxyamine and a compound having a structure of Formula I:



Formula I

20

wherein X is O or NH,

Y is O, S, or NH,

Z is absent or represents O, S, or NH, and

R represents a hydrogen or a hydrocarbon moiety,

and pharmaceutically acceptable salts thereof.

206. The kit of any one of claims 188 or 189, wherein said BER inhibitor is a PARP inhibitor.

207. The kit of claim 206, wherein said PARP inhibitor is selected from
5 PD128763, 3-AB, and 6-AN.

208. The kit of claim 206, wherein said kit further comprises a topoisomerase inhibitor.

209. The kit of claim 208, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.

10 210. The kit of claim 212, wherein the topoisomerase II inhibitor is etoposide.

211. The kit of claim 172, wherein the amount of the anticancer agent is subtherapeutic when administered in the absence of the base excision repair inhibitor.

15 212. The kit of claim 172, wherein said base excision repair inhibitor is selected from an AP endonuclease inhibitor, a DNA glycosylase inhibitor, a DNA polymerase inhibitor, a PARP inhibitor, and a DNA ligase inhibitor.

213. The kit of claim 212, wherein said base excision repair inhibitor is an AP endonuclease inhibitor.

20 214. The kit of claim 212, wherein said base excision repair inhibitor is a PARP inhibitor.

215. The kit of claim 214, wherein said PARP inhibitor is selected from PD128763, 3-AB, and 6-AN.

216. The kit of claim 212, wherein said BER inhibitor is an inhibitor of DNA polymerase.

217. The kit of claim 216, wherein said inhibitor of DNA polymerase inhibits DNA polymerase β , γ , or ϵ .
218. The kit of claim 212, wherein said base excision repair inhibitor is a DNA ligase inhibitor.
- 5 219. The kit of claim 218, wherein said DNA ligase inhibitor inhibits the action of DNA ligase I or DNA ligase II.
220. The kit of any one of claims 178, 181, 184, 187, 191, 195 or 203, wherein the instructions are for administering the kit to a patient diagnosed with cancer.
221. The kit of claim 220, wherein the instructions are for administering the
10 pharmaceutical preparation orally.
222. The kit of claim 220, wherein the instructions are for administering the pharmaceutical preparation intravenously.
223. The kit of claim 220, wherein the cancer is selected from carcinomas, melanomas, sarcomas, lymphomas, leukemias, astrocytomas, gliomas, malignant
15 melanomas, chronic lymphocytic leukemia, lung cancers, and breast cancers.
224. The kit of claim 172, wherein said kit further comprises a DNA alkyltransferase inhibitor.
225. The kit of claim 224, wherein said DNA alkyltransferase inhibitor is BG.
226. The kit of claim 172, wherein said kit further comprises a topoisomerase
20 inhibitor.
227. The kit of claim 226, wherein said topoisomerase inhibitor is a topoisomerase I inhibitor.
228. The kit of claim 226, wherein said topoisomerase inhibitor is a topoisomerase II inhibitor.
- 25 229. The kit of claim 228, wherein the topoisomerase II inhibitor is etoposide.

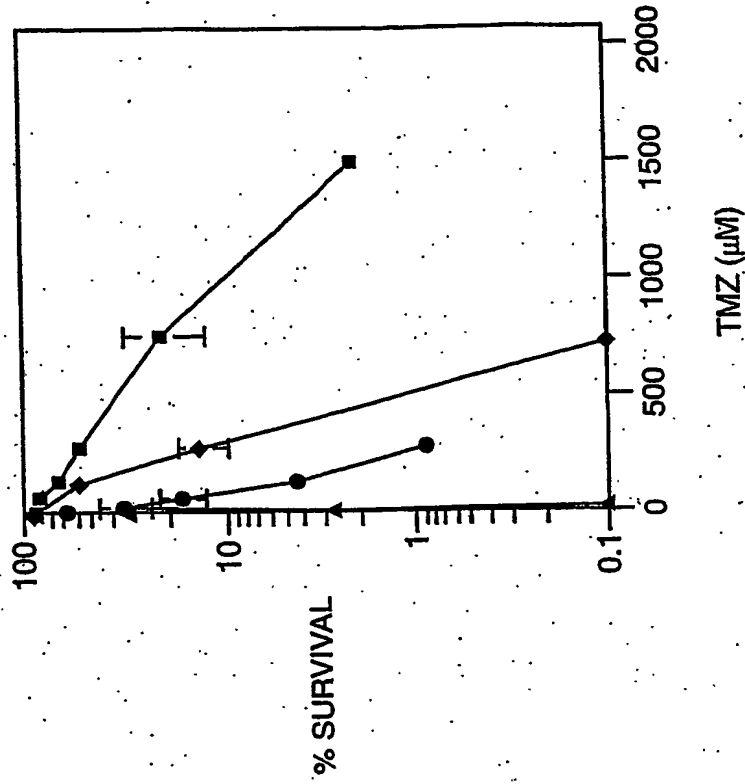


FIG. 1B

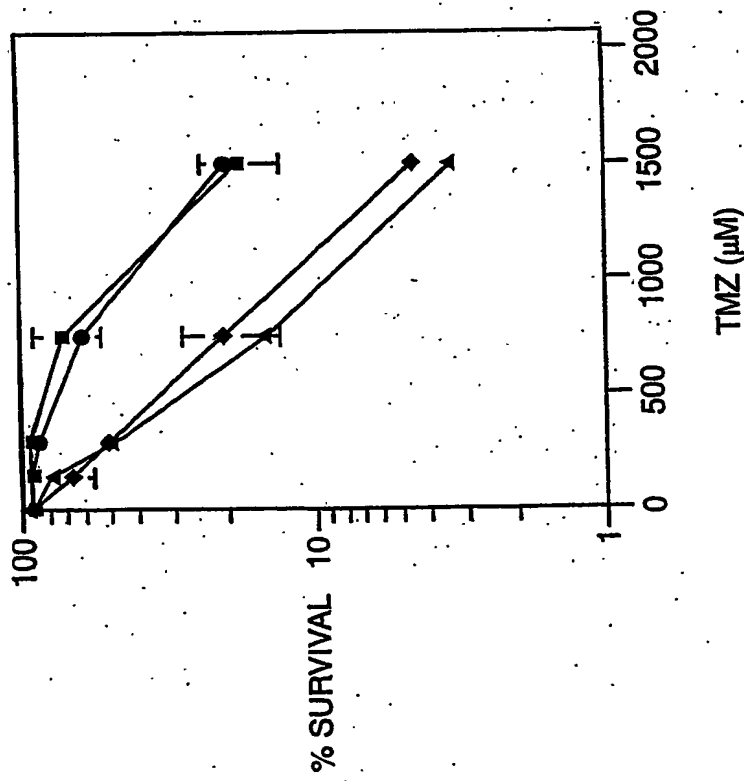


FIG. 1A

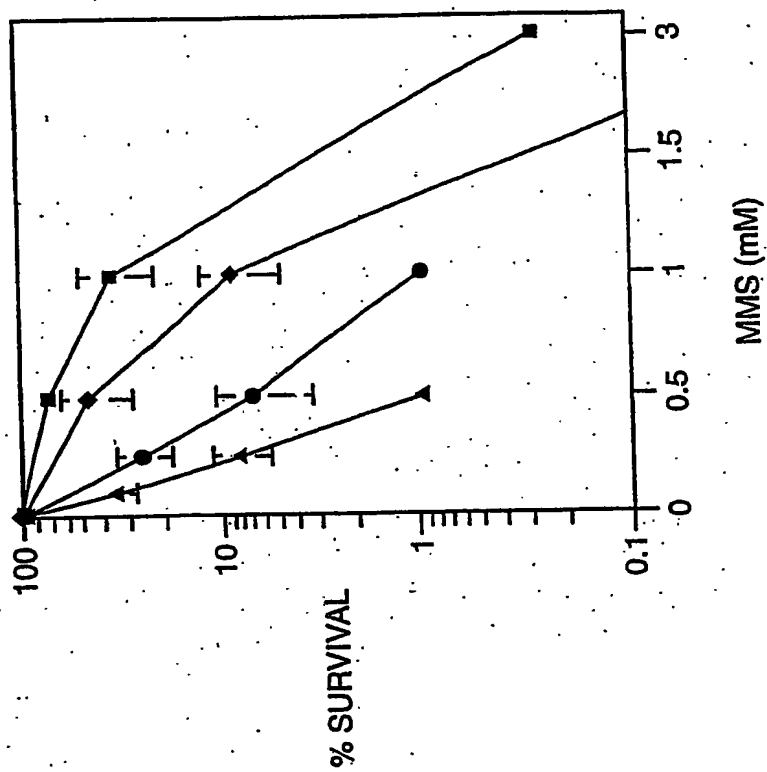


FIG. 2B

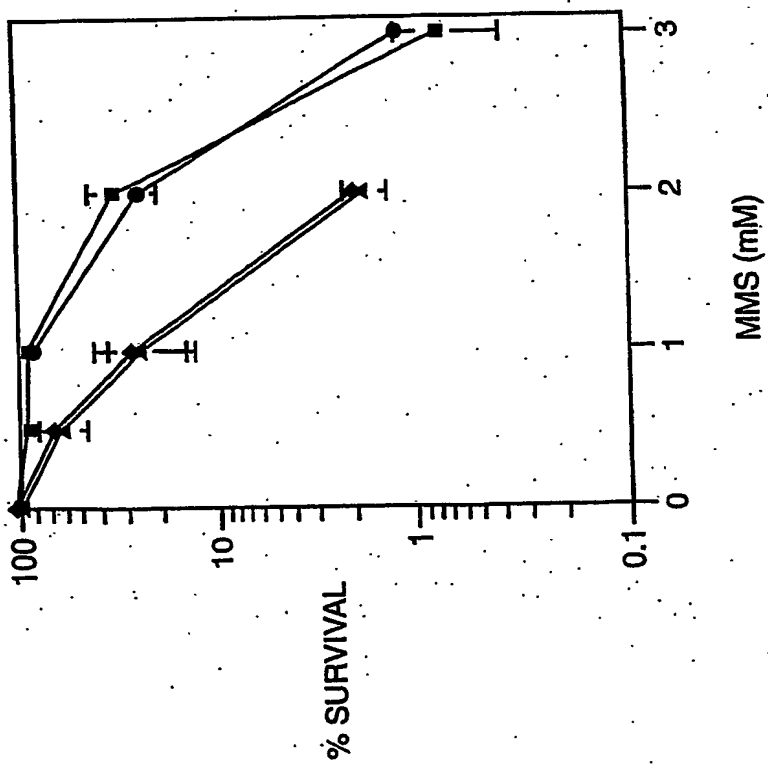


FIG. 2A

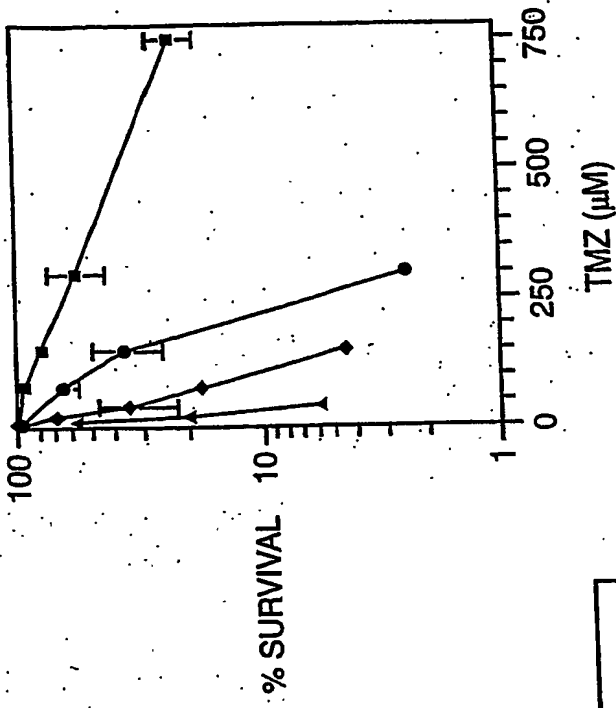


FIG. 3A

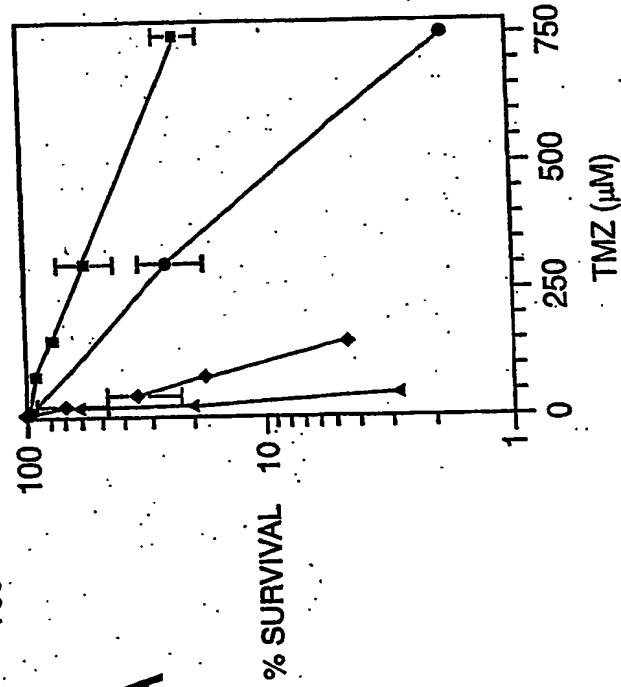


FIG. 3B

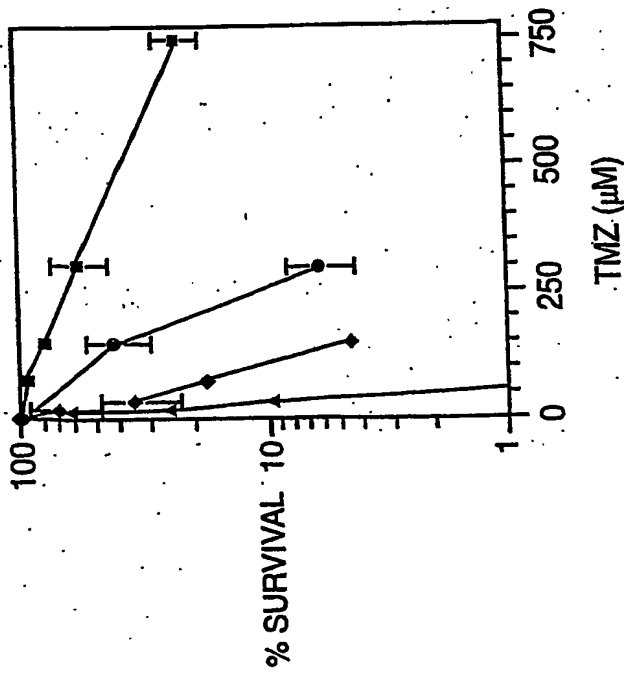


FIG. 3C

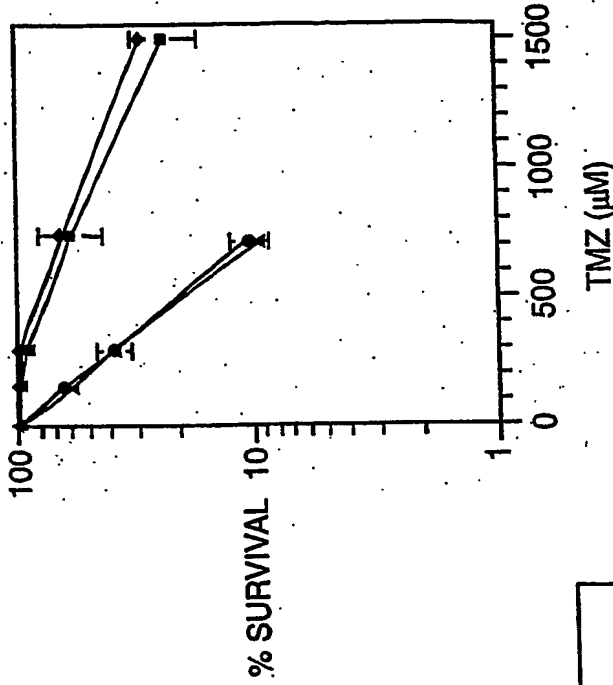


FIG. 4B

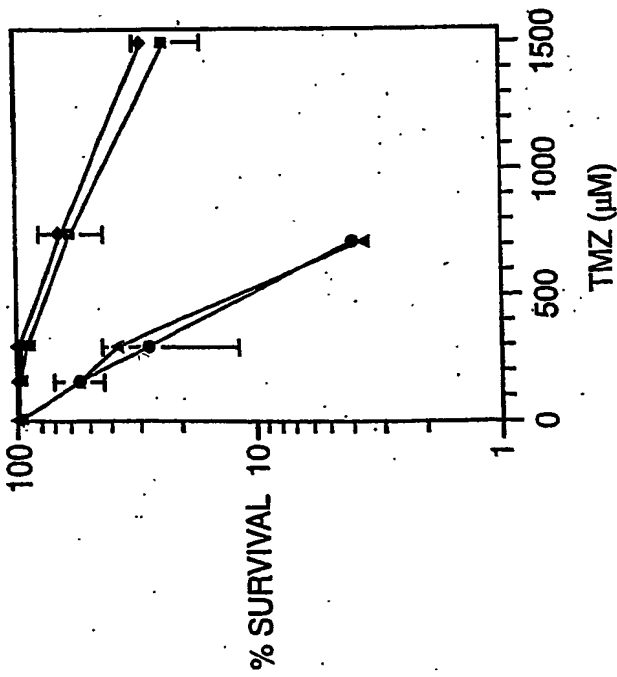


FIG. 4A

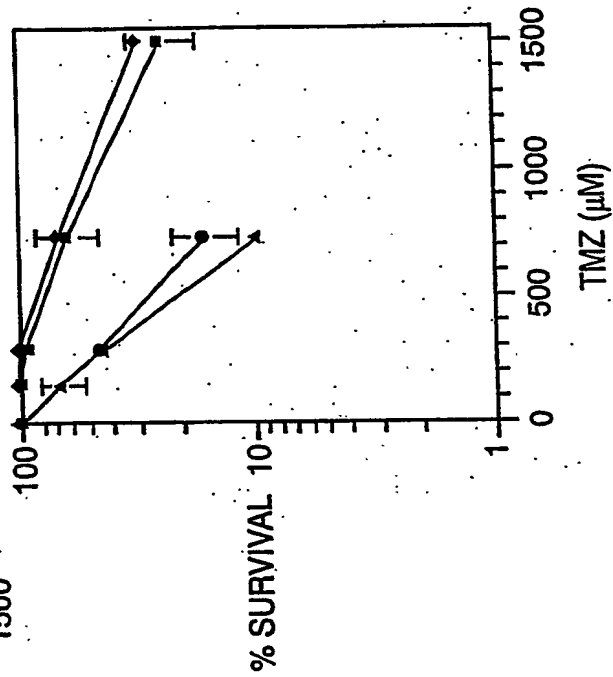


FIG. 4C

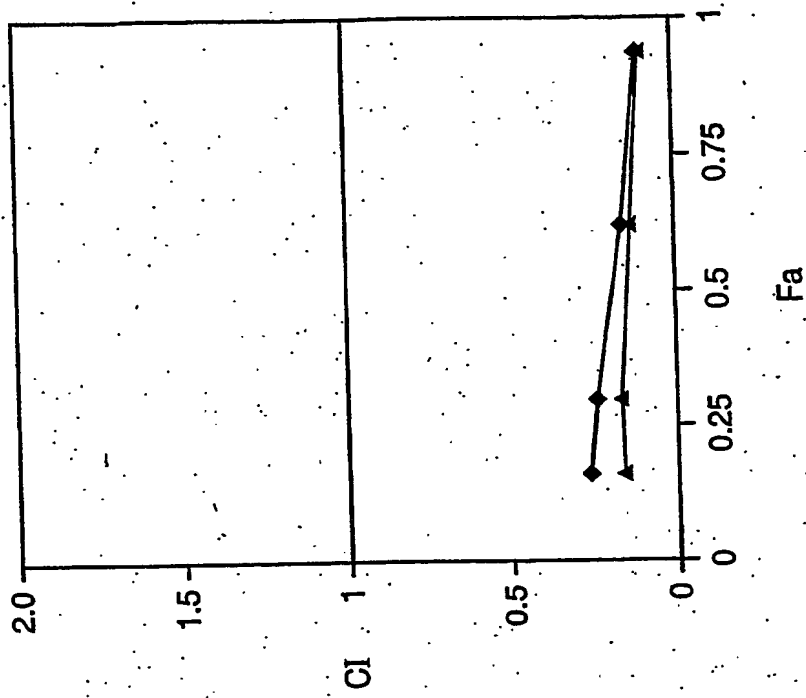


FIG. 5A

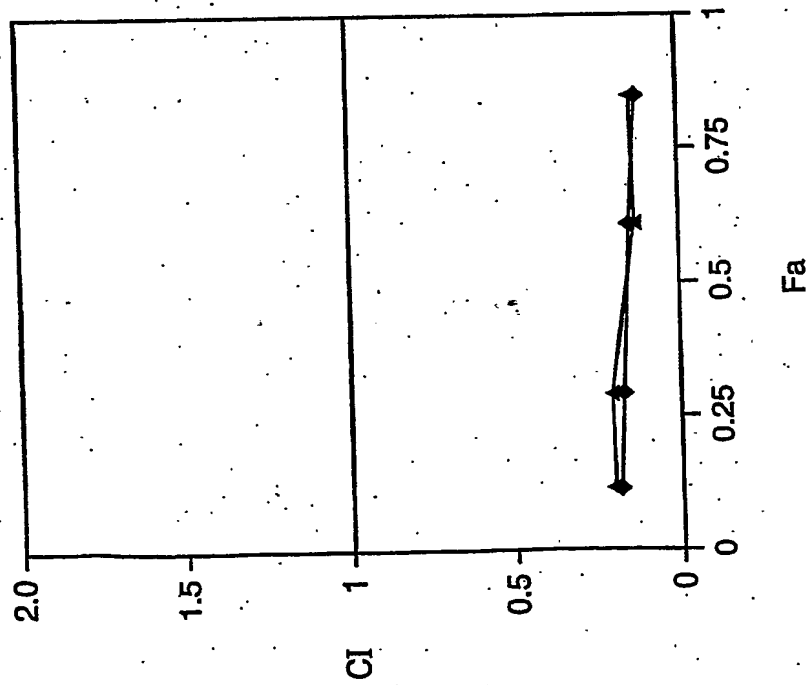


FIG. 5B

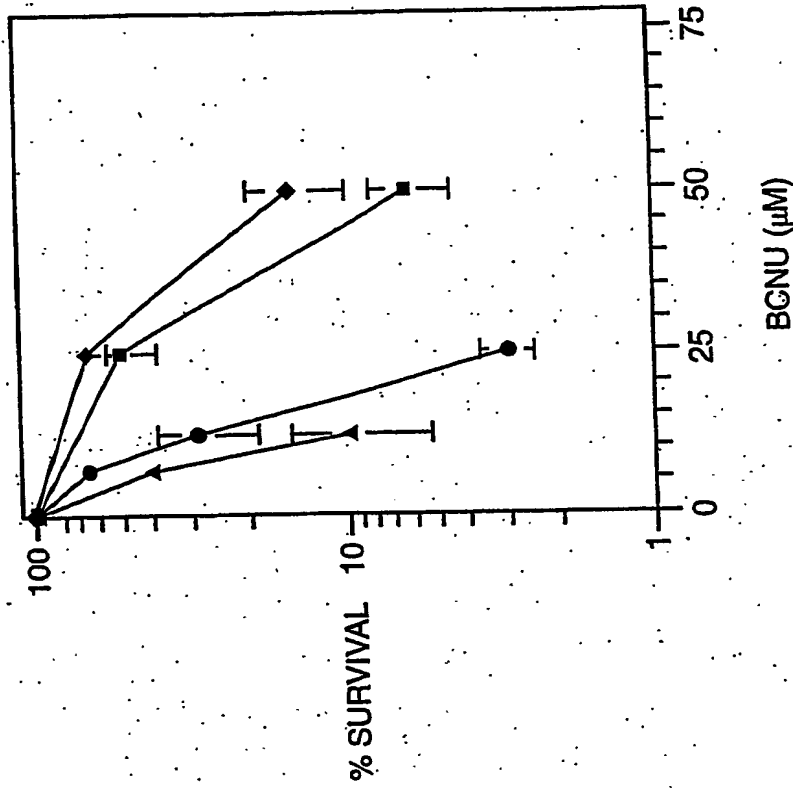


FIG. 6B

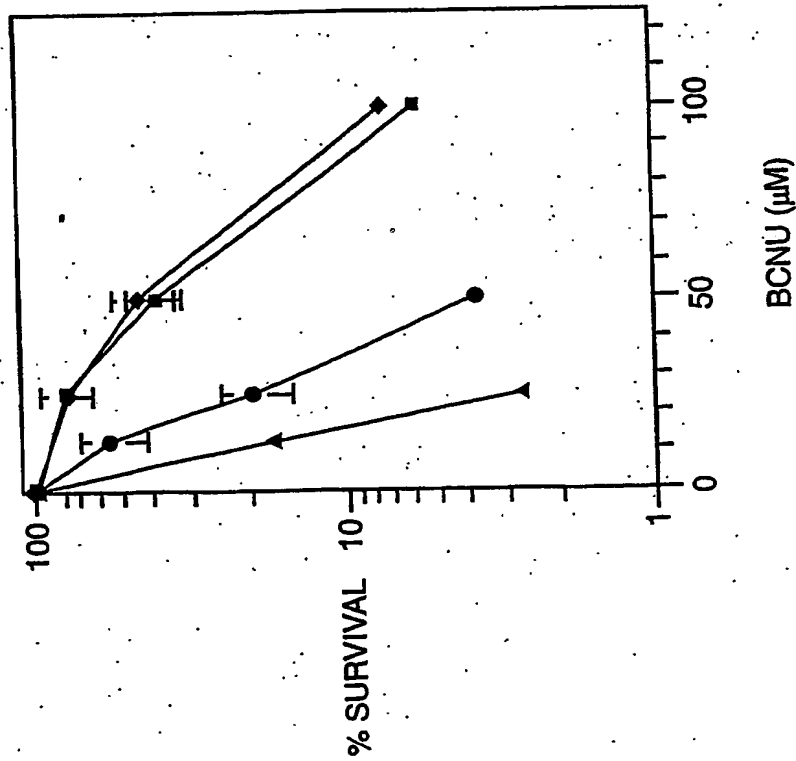


FIG. 6A

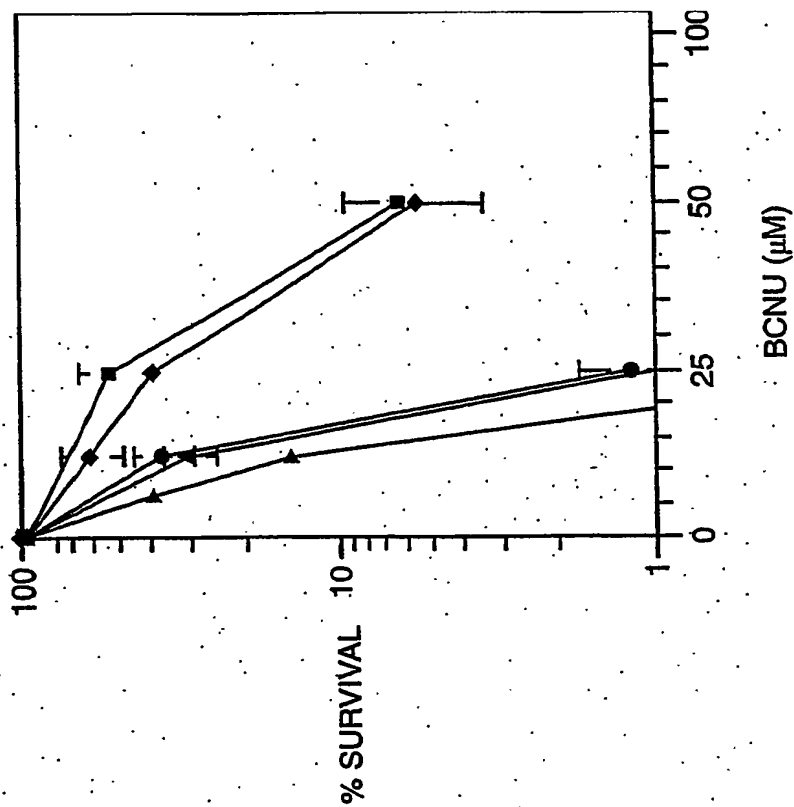


FIG. 7B

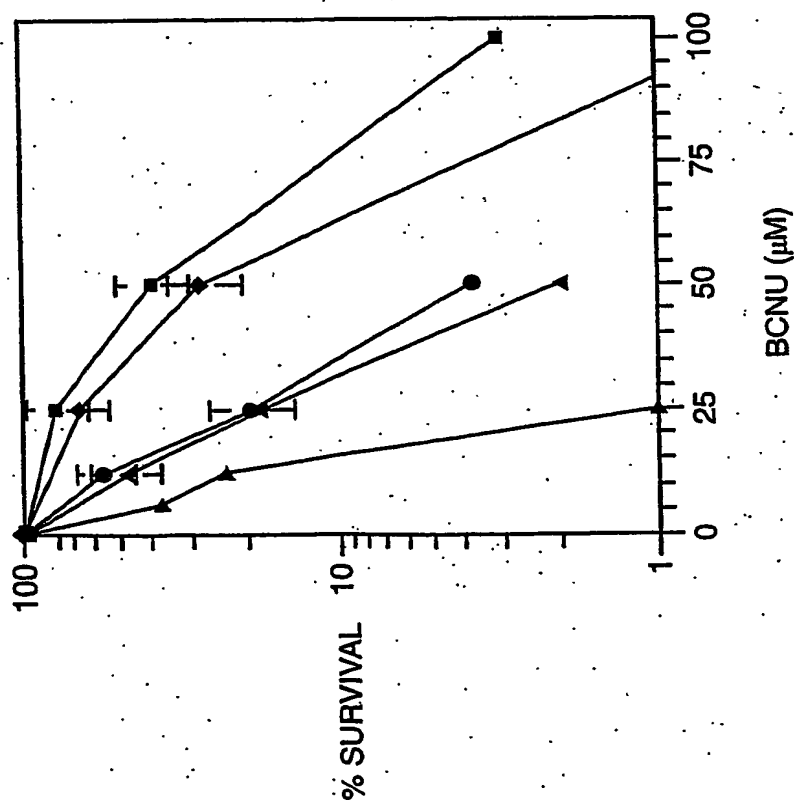


FIG. 7A

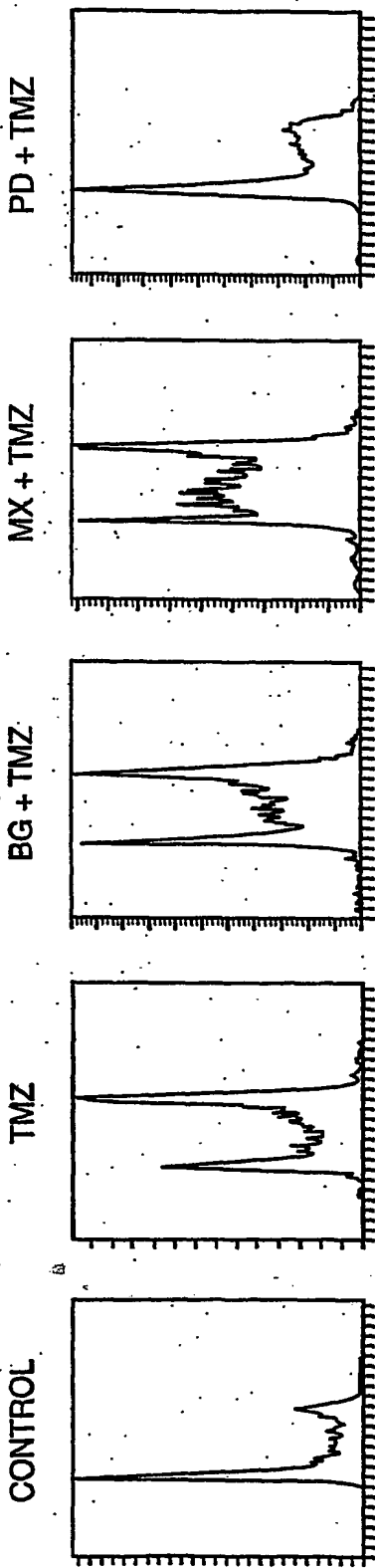


FIG. 8A

FIG. 8B

FIG. 8C

FIG. 8D

FIG. 8E

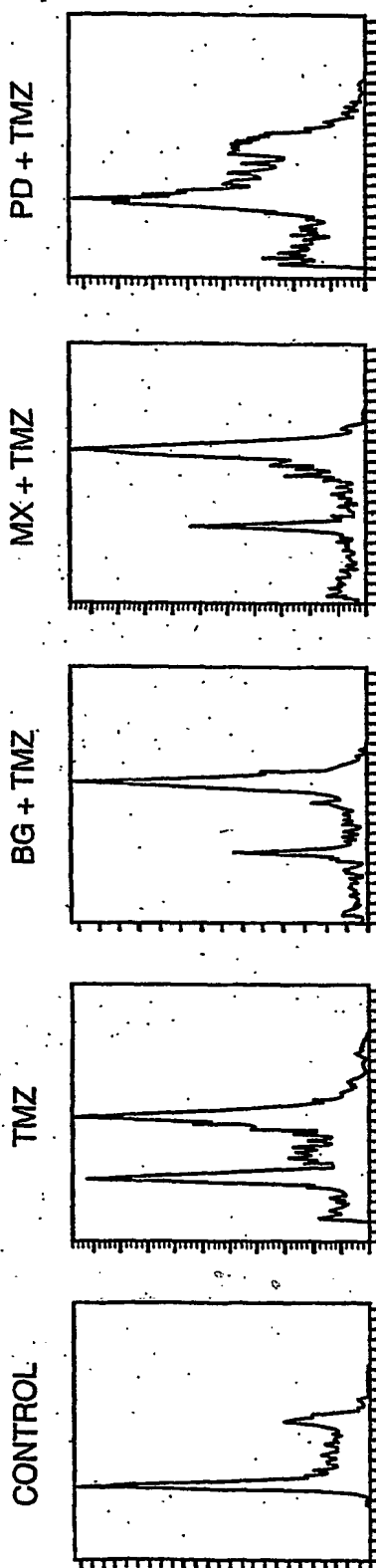


FIG. 8F

FIG. 8G

FIG. 8H

FIG. 8I

FIG. 8J

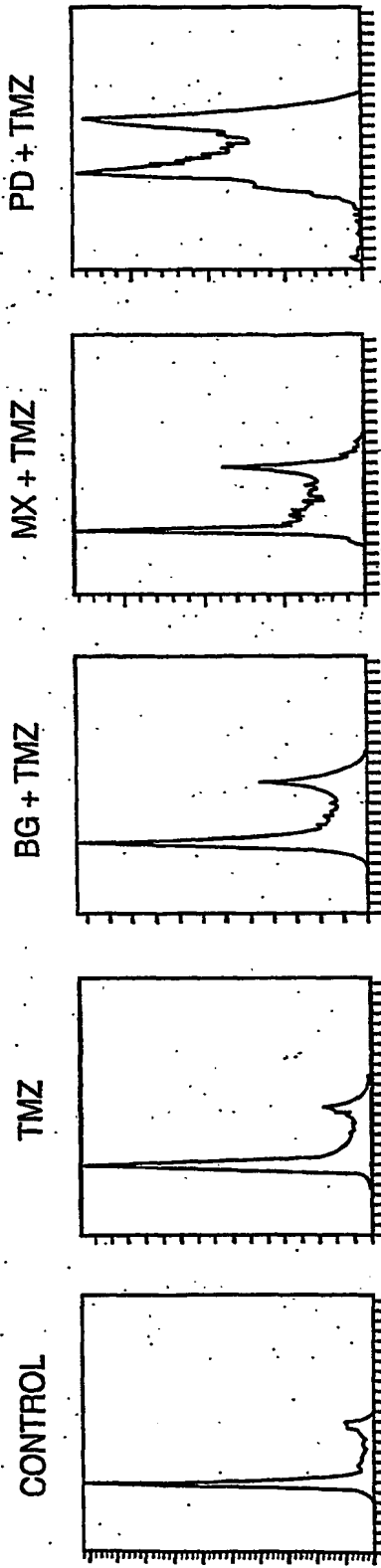


FIG. 8K FIG. 8L FIG. 8M FIG. 8N FIG. 8O

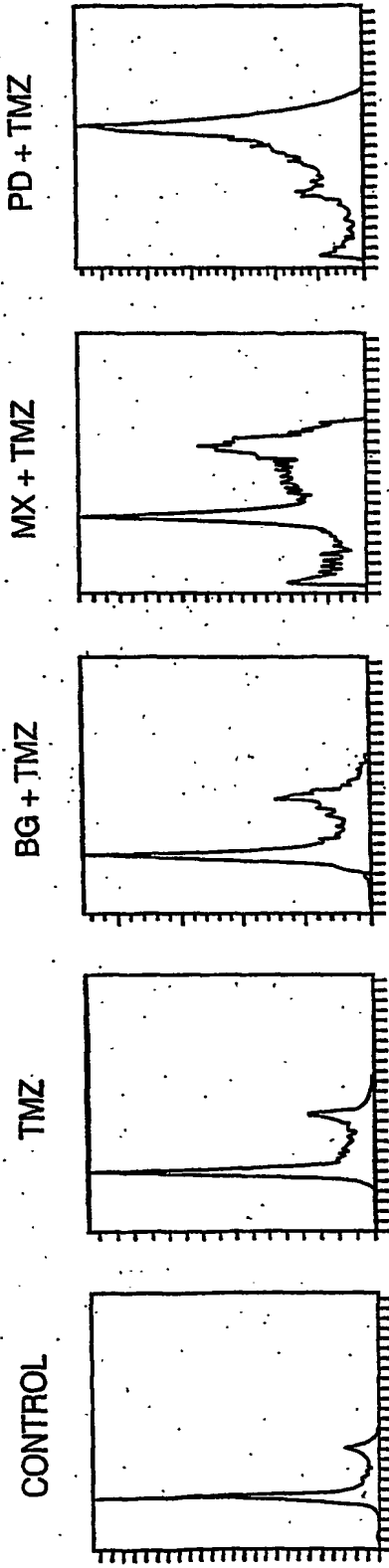


FIG. 8P FIG. 8Q FIG. 8R FIG. 8S FIG. 8T

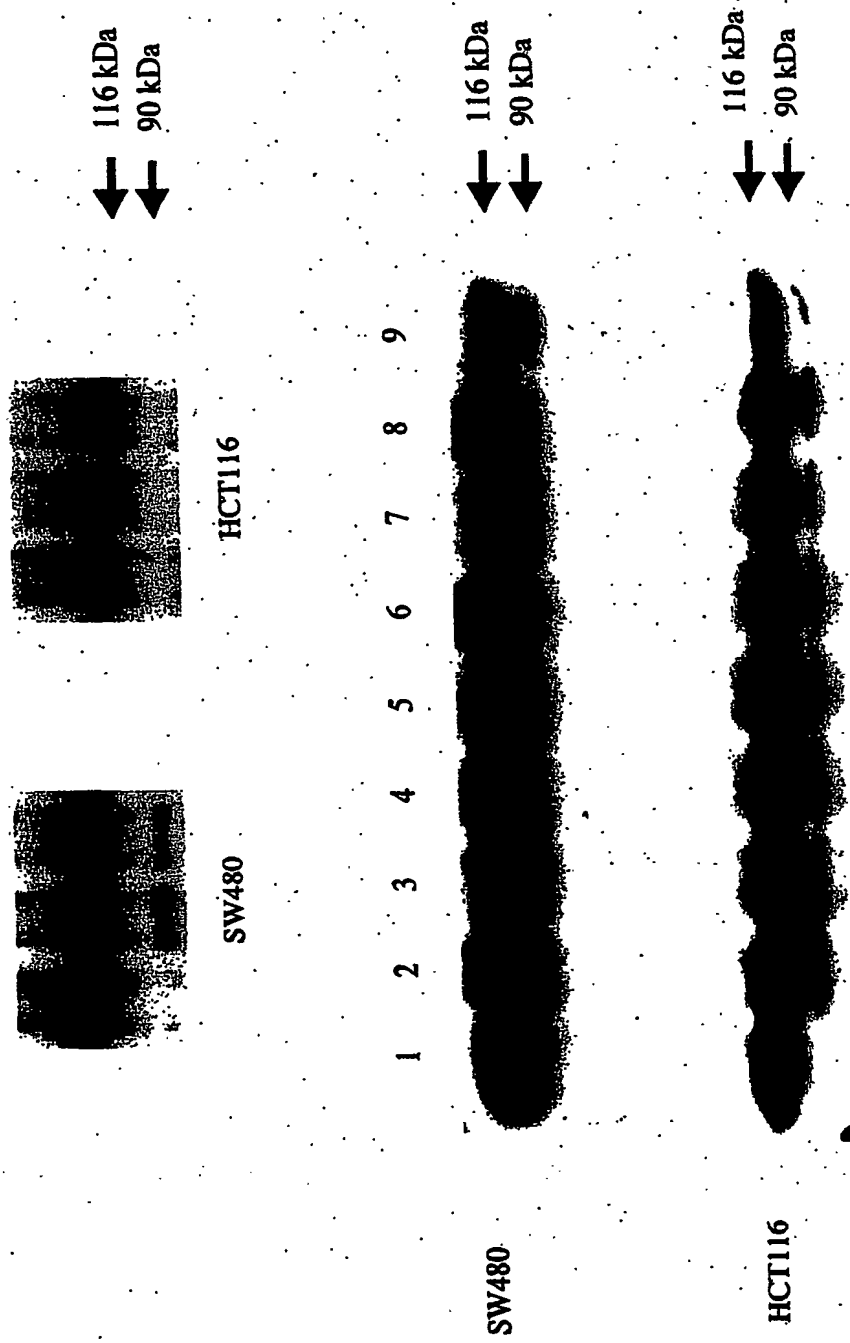


FIG. 9

11/26

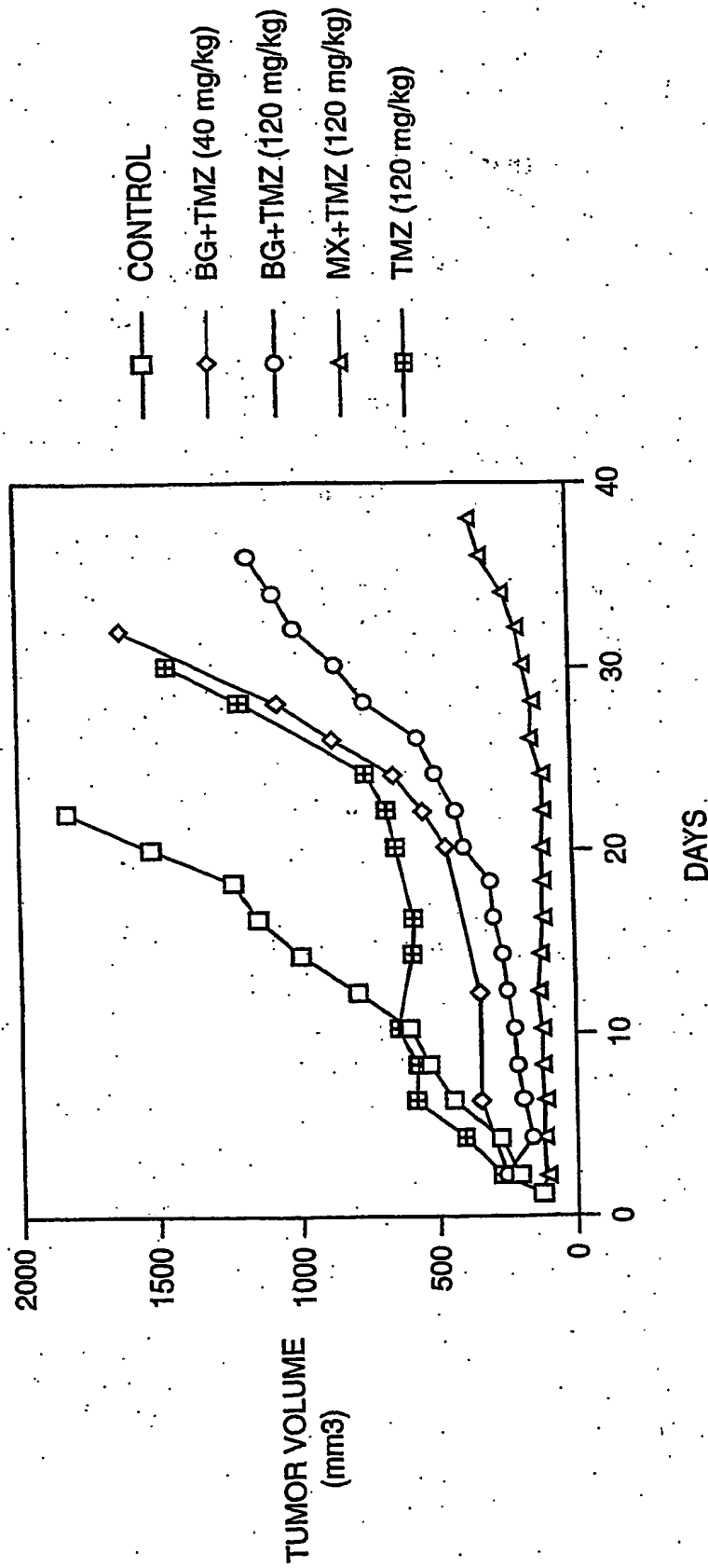


FIG. 10

12/26

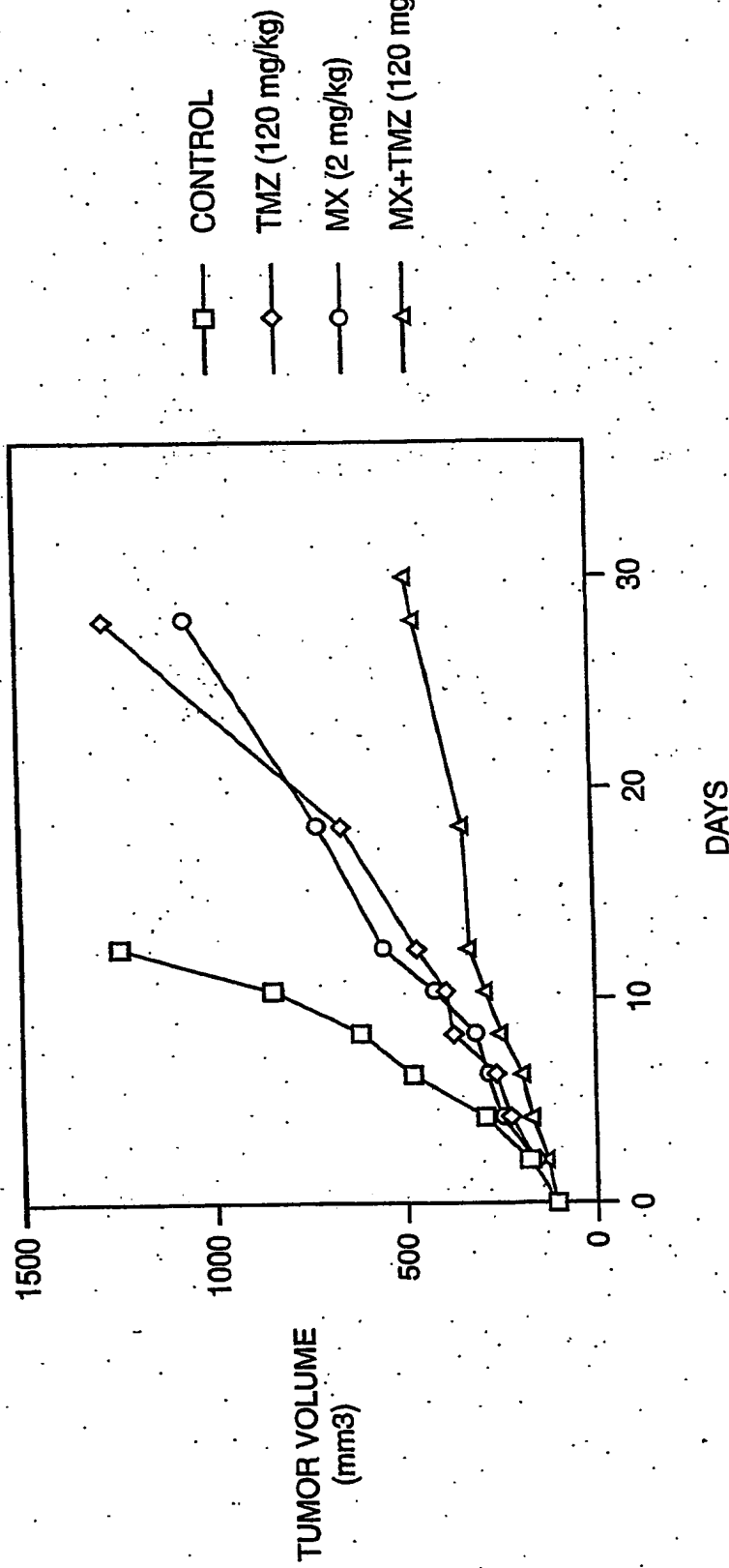


FIG. 11

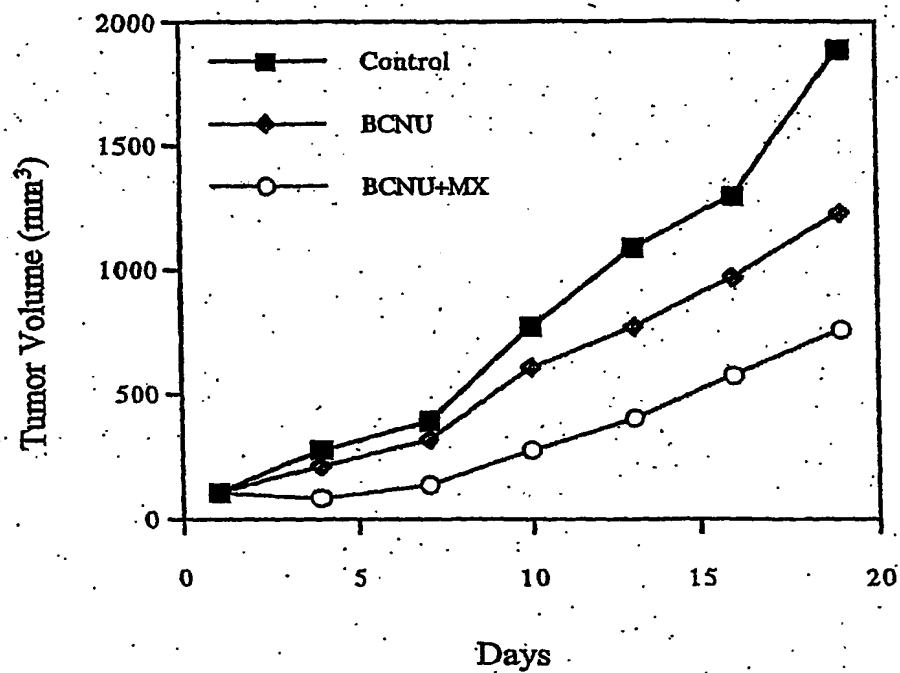


FIG. 12A

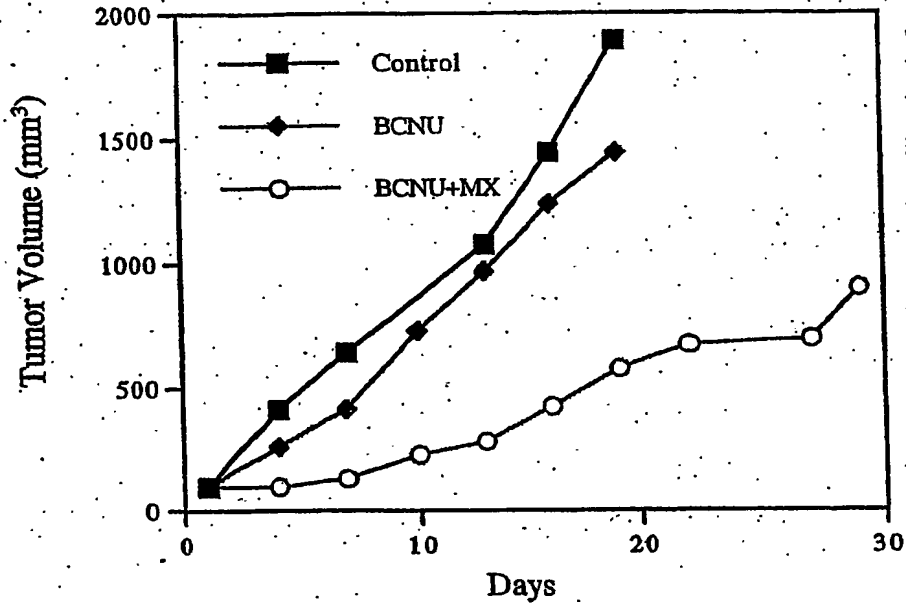


FIG. 12B

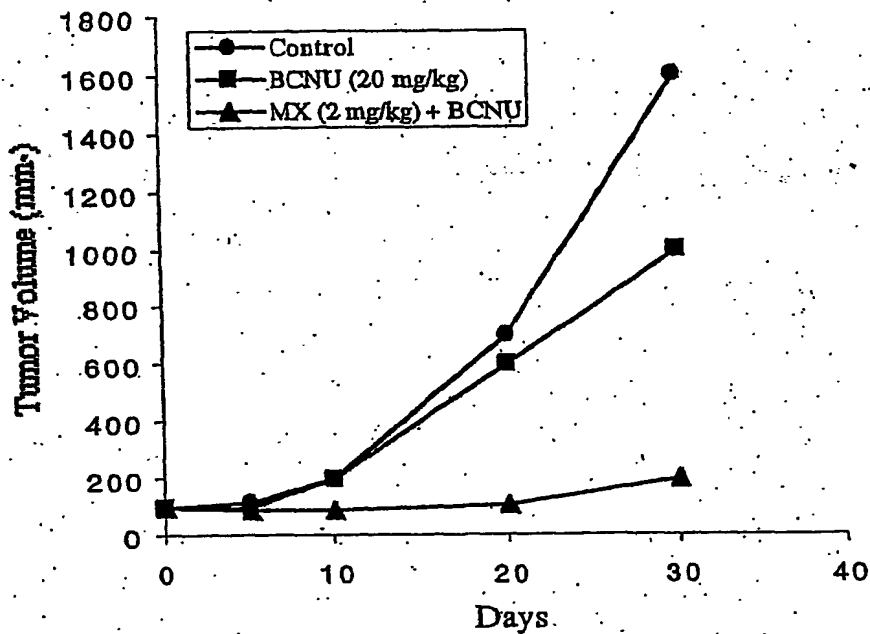


FIG. 12C

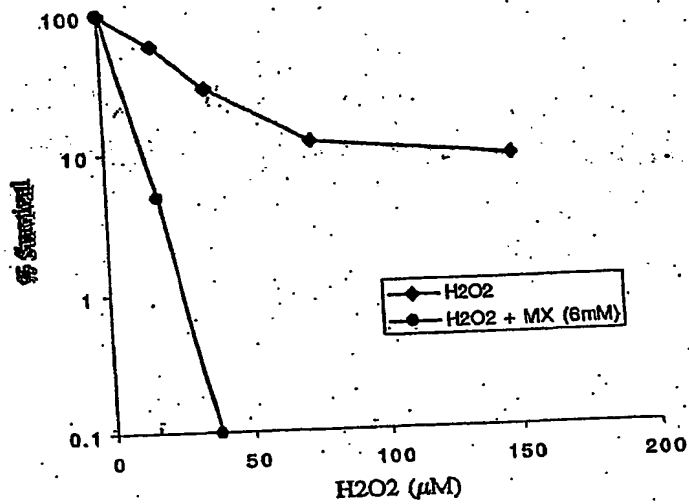
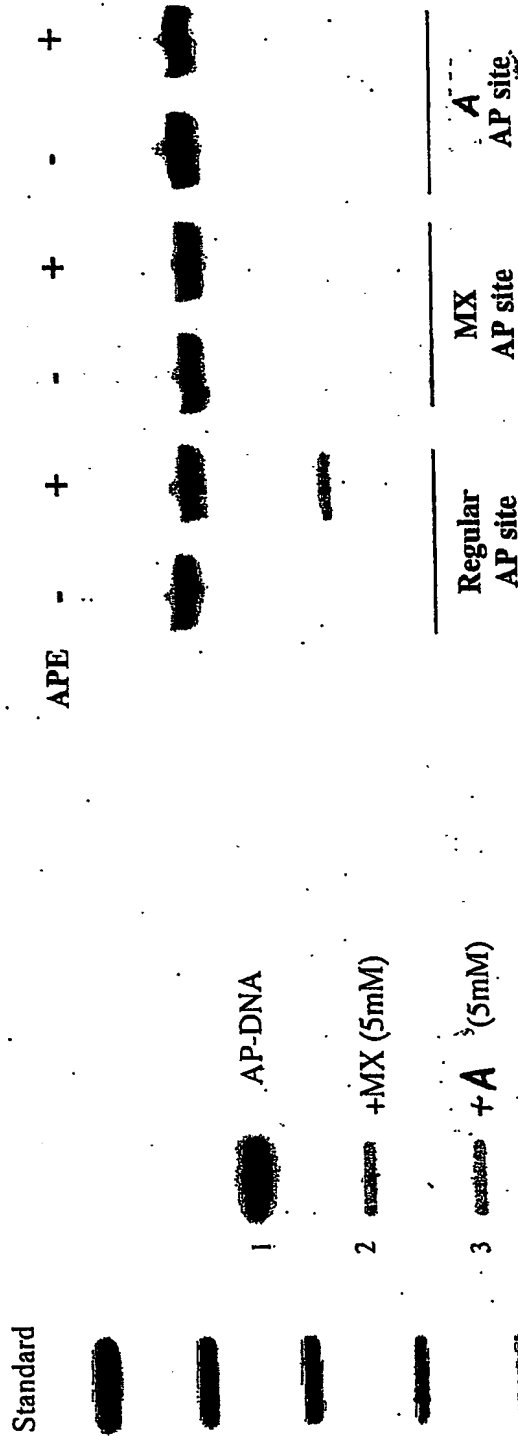


FIG. 13

Fig 1A

b. APE cleavage in regular and modified AP site



a. AP-site assay by ARP

A and MX have the same reactivity to AP sites. Results show that these two compounds bind to AP sites (A) and result in resistance to cleavage of APE (B), an initial step of repair AP site in base excision repair pathway.

Fig 15

Potential of TMZ-Cytotoxicity by A in Tumor cells

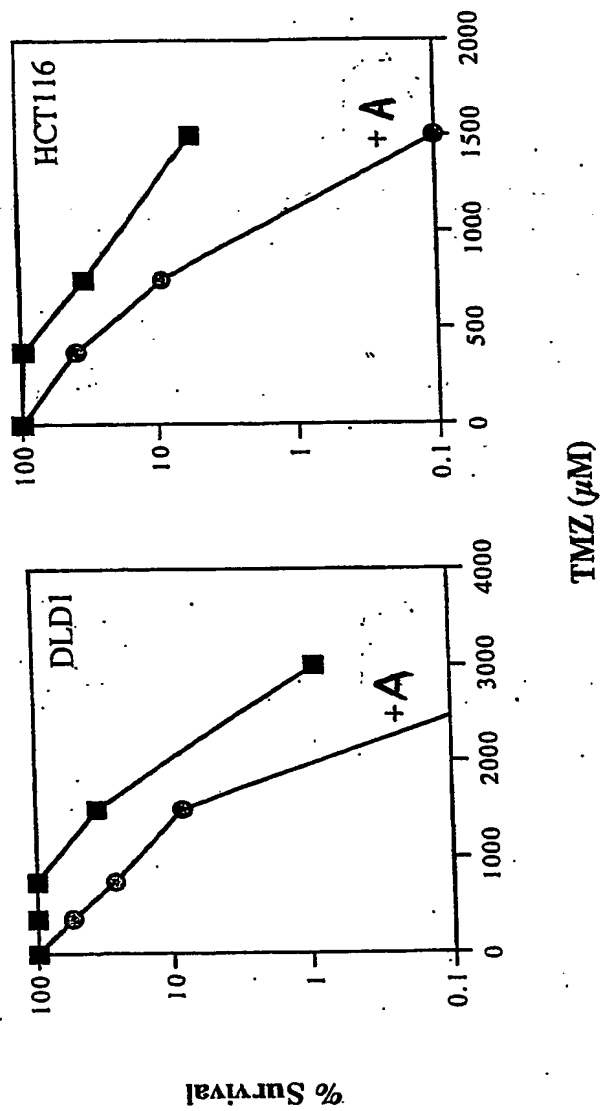
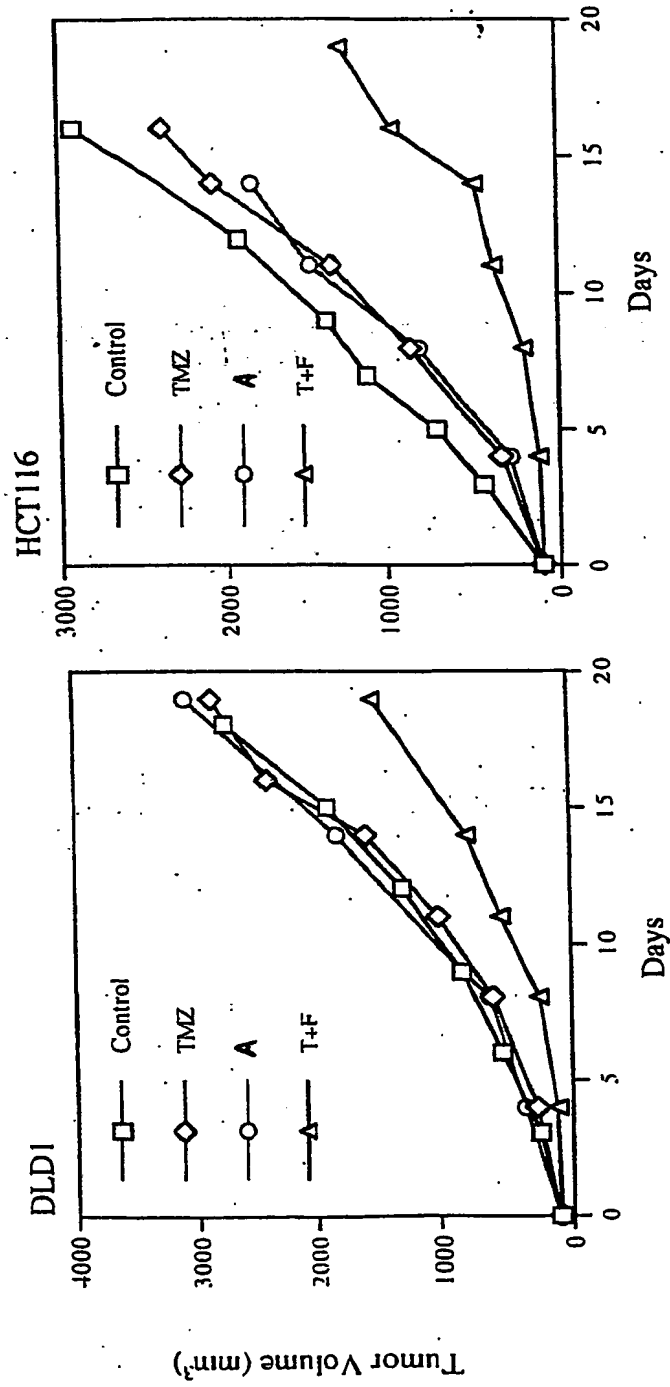


Fig. 16

Inhibition of Tumor Growth by TMZ plus A in Nude Mice



Schematic diagram of the Preparation of DNA substrates with regular AP site or MX-AP sites

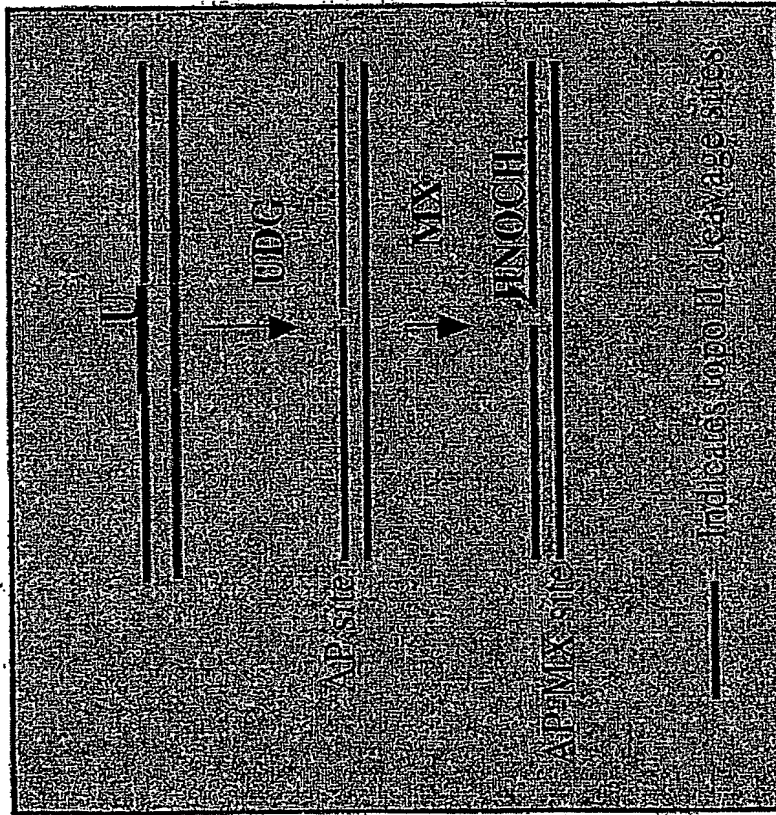


Figure 17

MX bound AP Site is Resistant to Cleavage by AP-endonuclease

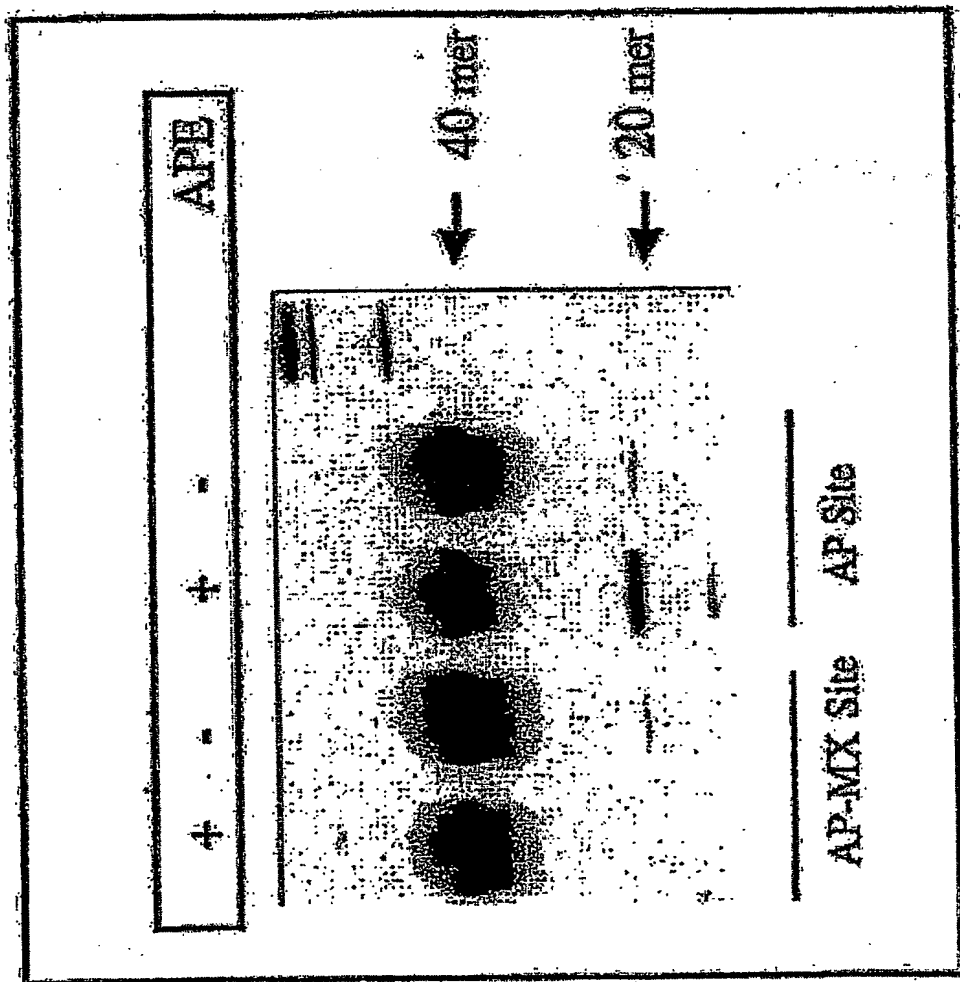
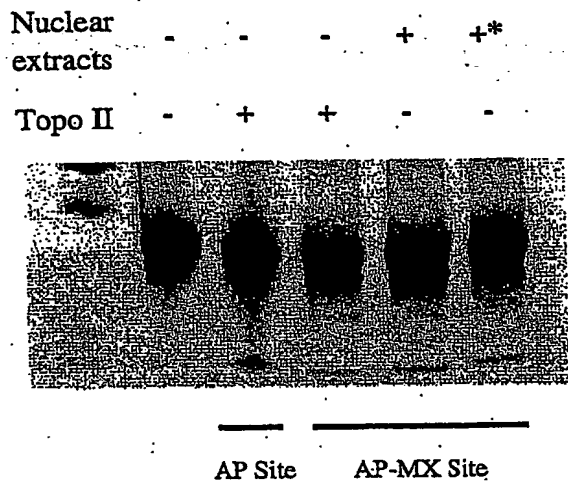


Figure 18

Figure 14

Topo II-mediated DNA Cleavage in Oligonucleotides Containing a Topo II Cleavage Site



* SW480 and HCT116 cells were treated with etoposide(5 μ M) for 4 hrs

Figure 20

Comparison of DNA Double Strand Breaks Induced by The Combination of TMZ, MX and Etoposide

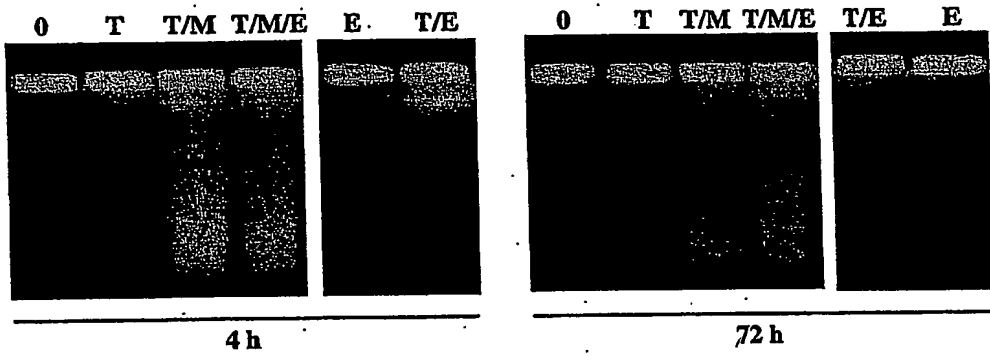


Figure 21

Synergy Analysis of The Interaction Between TMZ/MX and Etoposide in Cell Line SW480

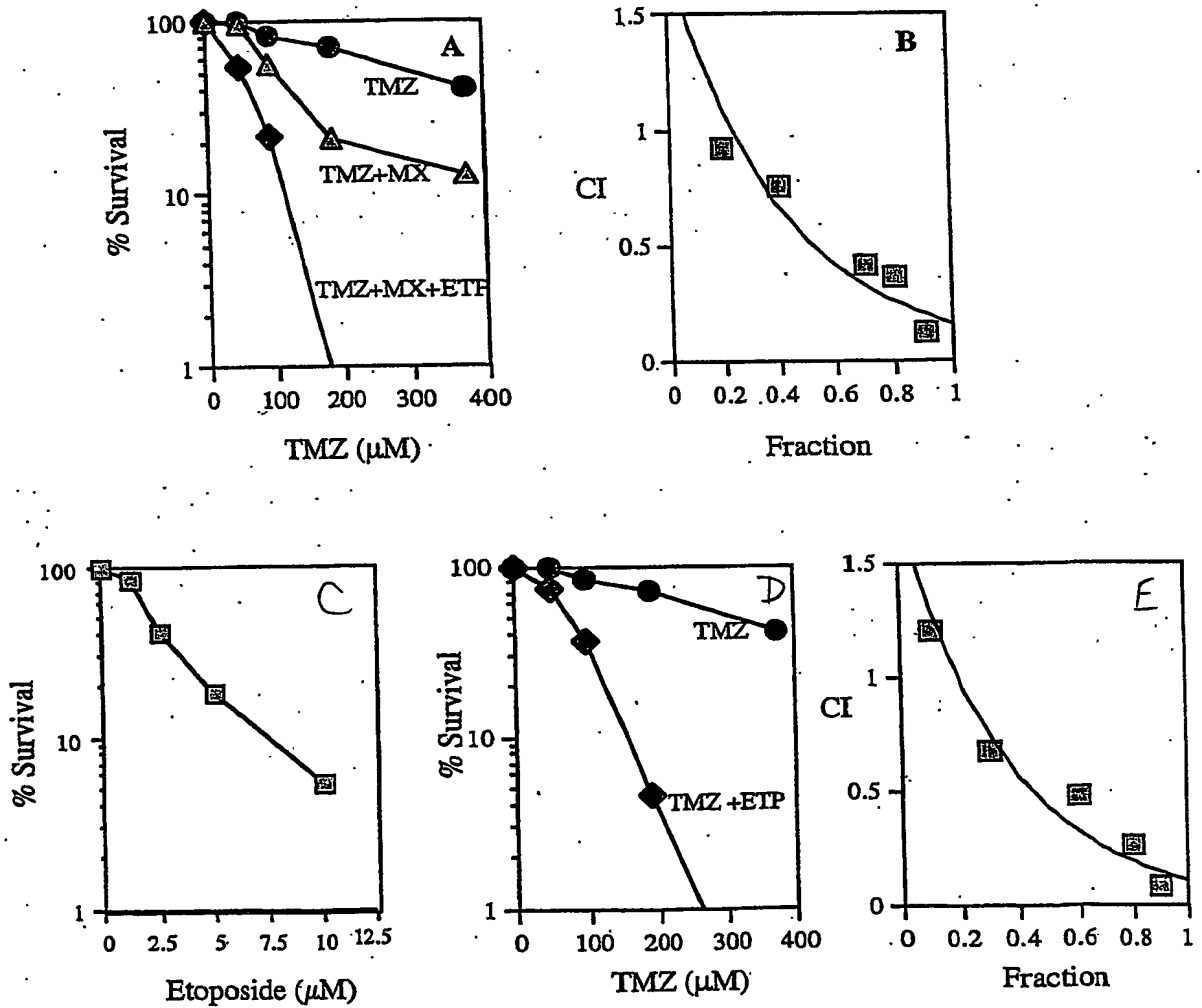


Fig 22

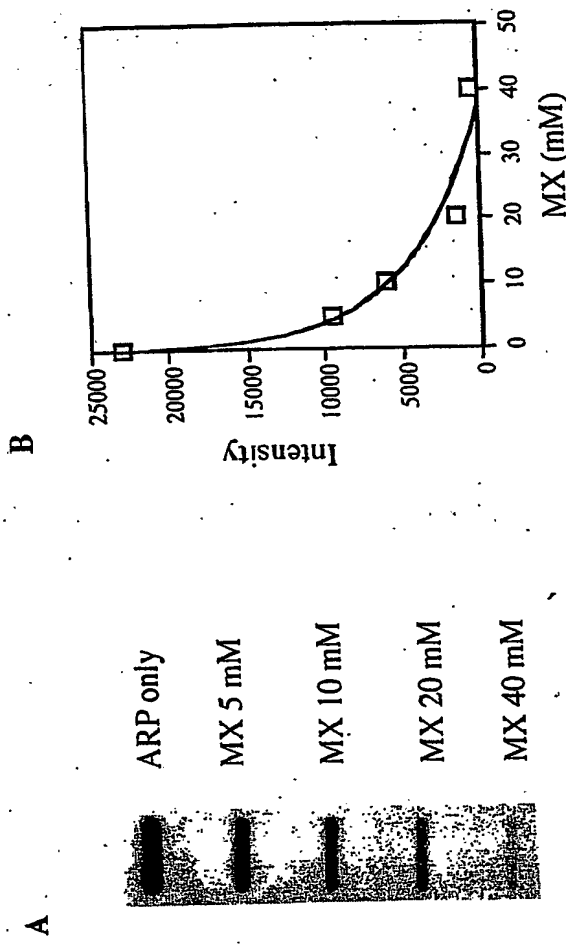


Fig 22A: The images of ARP-AP sites showed that MX inhibited detection of AP sites by ARP, resulting from competitive reactivity with AP sites between MX and ARP.

Fig 22B: Relative reduction of AP sites estimated by densitometric intensity showed that MX (5 – 20 mM) blocked 38% to 80% AP binding sites.

Fig 23

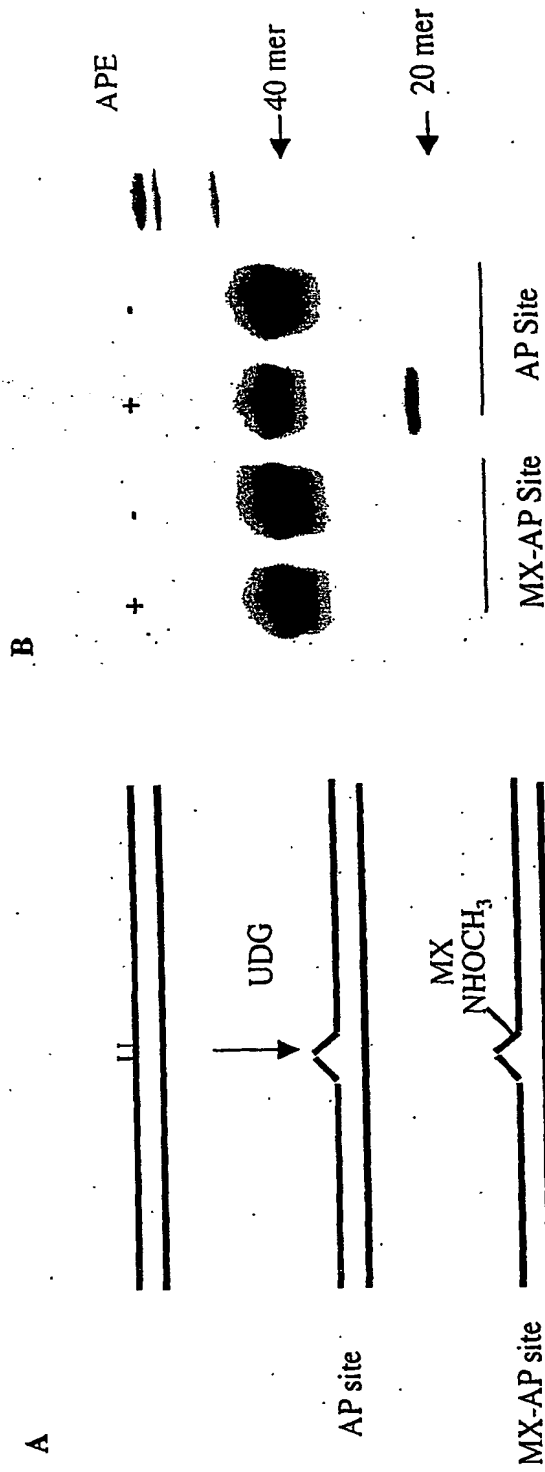


Fig 23 A: A diagram for preparation of DNA substrates containing either regular AP sites or MX-modified AP sites

Fig 23 B: MX-AP site is resistant to the cleavage by APE

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US03/05032

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) :A61K 31/13, 31/33

US CL :Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/34, 50, 51, 183, 233.5, 262, 263, 264, 299, 309, 449, 588, 589, 599, 644, 645, 665, 666

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 6,197,785 B1 (JACKSON ET AL) 06 March 2001 (06.03.01), see the entire document, especially column 54, line 7 to column 55, line 40).	1-229

Further documents are listed in the continuation of Box C. See patent family annex.

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"B" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
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"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 01 JUNE 2003	Date of mailing of the international search report 18 JUN 2003
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Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer FREDERICK BRASS Telephone No. (703) 308-1235
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US03/05032

A. CLASSIFICATION OF SUBJECT MATTER:
US CL :

514/34, 50, 51, 183, 233.5, 262, 263, 264, 299, 309, 449, 588, 589, 599, 644, 645, 665, 666

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